

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61K 7/06, 7/13		A2	(11) International Publication Number: WO 99/13823 (43) International Publication Date: 25 March 1999 (25.03.99)
(21) International Application Number: PCT/IB98/01381		(74) Agents: REED, T., David et al.; The Procter & Gamble Company, 5299 Spring Grove Avenue, Cincinnati, OH 45217-1087 (US).	
(22) International Filing Date: 4 September 1998 (04.09.98)			
(30) Priority Data: PCT/US97/16412 17 September 1997 (17.09.97) US PCT/US97/16617 17 September 1997 (17.09.97) US PCT/US97/16618 17 September 1997 (17.09.97) US PCT/US97/16619 17 September 1997 (17.09.97) US PCT/US97/16615 17 September 1997 (17.09.97) US		(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).	
(71) Applicant (for all designated States except US): THE PROCTER & GAMBLE COMPANY [US/US]; One Procter & Gamble Plaza, Cincinnati, OH 45202 (US).		Published <i>Without international search report and to be republished upon receipt of that report.</i>	
(72) Inventors; and (75) Inventors/Applicants (for US only): MITSUMATSU, Arata [JP/JP]; 6-16-306, Midorigaoka-cho, Ashiya-shi, Hyogo 659 (JP). LUO, Xiaochun [JP/JP]; 5-15-703, Koyo-cho Naka, Higashinada-ku, Kobe 658 (JP). UCHIYAMA, Hiro-taka [JP/JP]; 5-8-17-3B, Nishi-Okamoto, Higashinada-ku, Kobe 658 (JP). SETSER, Julia, Leet [US/JP]; 2101, 5-15, Koyo-cho Naka, Higashinada-ku, Kobe 658 (JP).			

(54) Title: HAIR CARE COMPOSITIONS COMPRISING OPTICAL BRIGHTENERS AND HAIR CONDITIONING AGENTS

(57) Abstract

Disclosed are hair care compositions comprising: (a) an effective amount of an optical brightener; and (b) a hair conditioning agent selected from the group consisting of silicone compounds, cationic compounds, high melting point compounds, perfume compounds, water-insoluble high molecular weight oily compounds, and mixtures thereof.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon			PT	Portugal		
CN	China	KR	Republic of Korea	RO	Romania		
CU	Cuba	KZ	Kazakhstan	RU	Russian Federation		
CZ	Czech Republic	LC	Saint Lucia	SD	Sweden		
DE	Germany	LI	Liechtenstein	SE	Sri Lanka		
DK	Denmark	LK	Sri Lanka	SG	Singapore		
EE	Estonia	LR	Liberia				

HAIR CARE COMPOSITIONS COMPRISING OPTICAL BRIGHTENERS AND HAIR CONDITIONING AGENTS

TECHNICAL FIELD

The present invention relates to a hair care composition comprising optical brighteners which alter the color of the hair, while enhancing the shininess of the hair, and protecting the hair from further damage. More specifically, the present invention relates to a hair care composition comprising optical brighteners and hair conditioning agents.

BACKGROUND

The desire to regain the natural color and shine of damaged hair and the desire to alter the color of the hair to be more appealing are widely held. Damaged hair is perceived by the consumer as unfavorable appearances and less manageability of the hair. Such unfavorable appearances include alteration and fading of original color, less shine, and less luster.

A common way for alleviating the unfavorable appearances of damaged hair and to achieve appealing hair color is to dye the hair to the color desired. Dyeing the hair would provide the consumer with a stable color of hair for a relatively long period. However, dyeing the hair is generally time-consuming, cumbersome, and messy. Dyestuff may also be chemically harsh to the hair, scalp, and skin. The hair can be further damaged by dyeing. Thus, hair dye products are not suitable for daily use. Further, dyeing can leave the hair with a dull appearance, making the hair look less shiny.

Based on the foregoing, there is a need for a hair composition which can be used daily and which can alter the color of the hair, while enhancing the shininess of the hair, and protecting the hair from further damage.

Use of optical brighteners, or compounds otherwise described by names such as fluorescent whitening agents, fluorescent brighteners, or fluorescent

dyes, in the hair care field has been known in the art, such as in United States Patent 3,658,985, United States Patent 4,312,855, Canadian Patent 1,255,603, United States Patent 3,577,528, Great Britain Patent Specification 1,328,108, South African Application 676,049, European Publication 87,060, and Great Britain Patent Specification 2,307,639.

However, the use of optical brighteners alone may not provide sufficient conditioning benefits such as good combing, smoothness, and reduction of static. Thus there is also a need for a hair care compositions comprising optical brighteners and providing good conditioning benefits.

None of the existing art provides all of the advantages and benefits of the present invention.

SUMMARY

The present invention is directed to a hair care composition comprising: (a) an effective amount of an optical brightener; and (b) a hair conditioning agent selected from the group consisting of silicone compounds, cationic compounds, high melting point compounds, perfume compounds, water-insoluble high molecular weight oily compounds, and mixtures thereof.

These and other features, aspects, and advantages of the present invention will become evident to those skilled in the art from a reading of the present disclosure.

DETAILED DESCRIPTION

While the specification concludes with claims particularly pointing and distinctly claiming the invention, it is believed that the present invention will be better understood from the following description.

All percentages are by weight of the total composition unless otherwise indicated. All ratios are weight ratios unless otherwise indicated. All percentages, ratios, and levels of ingredients referred to herein are based on the actual amount of the ingredient, and do not include solvents, fillers, or other materials with which the ingredient may be combined as commercially available products, unless otherwise indicated.

As used herein, "comprising" means that other steps and other ingredients which do not affect the end result can be added. This term encompasses the terms "consisting of" and "consisting essentially of".

All cited references are incorporated herein by reference in their entireties. Citation of any reference is not an admission regarding any determination as to its availability as prior art to the claimed invention.

OPTICAL BRIGHTENERS

Optical brighteners are compounds which absorb ultraviolet light and re-emit the energy in the form of visible light. Specifically, the optical brighteners useful herein have an absorption, preferably a major absorption peak, between a wavelength of about 1nm and about 420nm, and an emission, preferably a major emission peak, between a wavelength of about 360nm and about 830nm; wherein the major absorption peak has a shorter wavelength than the major emission peak. More preferably, the optical brighteners useful herein have a major absorption peak between a wavelength of about 200nm and about 420nm, and a major emission peak between a wavelength of about 400nm and about 780nm. Optical brighteners may or may not have minor absorption peaks in the visible range between a wavelength of about 360nm and about 830nm. Optical brighteners can be described by other names in the art and in other industries, such as fluorescent whitening agents, fluorescent brighteners, and fluorescent dyes.

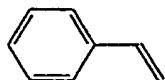
When applied to hair via suitable vehicles, optical brighteners herein provide benefits to the hair in three areas. First, optical brighteners herein alter the color of the hair by emitting light in the visible range. Second, optical brighteners herein enhance the shine of the hair by emitting light in the visible range. Third, optical brighteners herein protect the hair from ultraviolet light by absorbing ultraviolet light.

Optical brighteners in general are based on the structures of aromatic and heteroaromatic systems which provide these unique characteristics. The optical brighteners useful in the present invention can be classified according to their base structures, as described hereafter. Preferable optical brighteners herein include polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, and imidazoles, more preferably polystyrylstilbenes and hydrocoumarins.

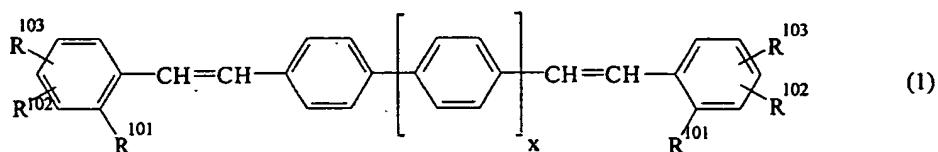
Preferably, optical brighteners herein are included in the hair care composition of the present invention at a level by weight of from about 0.001% to about 20%, more preferably from about 0.01% to about 10%.

Polystyrylstilbenes

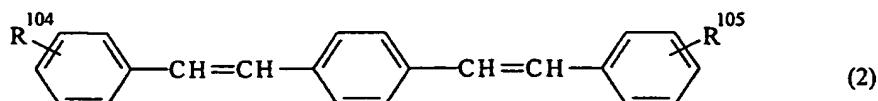
Polystyrylstilbenes are a class of compounds having two or more of the following base structure:



Polystyrylstilbenes useful in the present invention include those having formulae (1), (2) and (3):

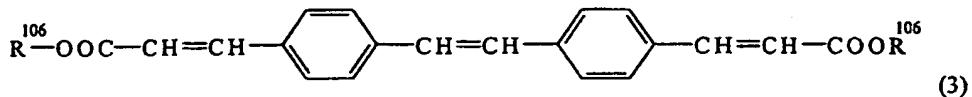


wherein R¹⁰¹ is H, OH, SO₃M, COOM, OSO₃M, OPO(OH)OM, wherein M is H, Na, K, Ca, Mg, ammonium, mono-, di-, tri- or tetra-C₁-C₃₀-alkylammonium, mono-, di- or tri-C₁-C₃₀-hydroxyalkylammonium or ammonium that is di- or tri-substituted with by a mixture of C₁-C₃₀-alkyl and C₁-C₃₀-hydroxyalkyl groups; or SO₂N(C₁-C₃₀-alkyl)₂, O-(C₁-C₃₀-alkyl), CN, Cl, COO(C₁-C₃₀-alkyl), CON(C₁-C₃₀-alkyl)₂ or O(CH₂)₃N⁺(CH₃)₂X⁻ wherein X⁻ is an anion of a chloride, bromide, iodide, formate, acetate, propionate, glycolate, lactate, acrylate, methanephosphonate, phosphite, dimethyl or diethyl phosphite anion; CN, or alkyl of 1 to 30 carbons, R¹⁰² and R¹⁰³, independently, are H, SO₃M wherein M is as previously defined; and x is 0 or 1; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably x is 1, R¹⁰¹ is SO₃Na and R¹⁰² and R¹⁰³ are H; wherein the compound has a trans-coplanar orientation;



wherein R¹⁰⁴ and R¹⁰⁵, independently, are CN, COO(C₁-C₃₀-alkyl), CONHC₁-C₄-alkyl, or CON(C₁-C₄-alkyl)₂, wherein the compound has a trans-coplanar

orientation or cis-coplanar orientation; preferably R¹⁰⁴ and R¹⁰⁵ is 2-cyano, wherein the compound has a trans-coplanar orientation; and



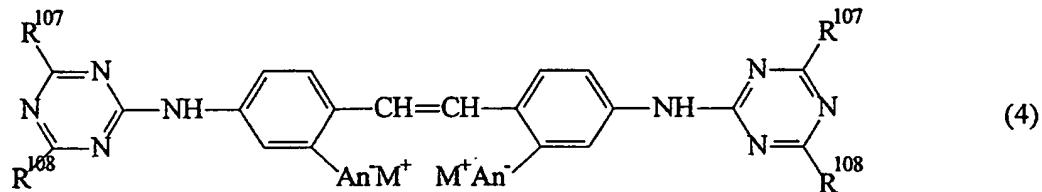
wherein each R¹⁰⁶, independently, is H, or alkyl of 1 to 30 carbons; and wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

Suitable polystyrylstilbenes include disodium-1, 4'-bis(2-sulfostyryl)bisphenyl (C.I. Fluorescent Brightener 351) with tradename Tinopal CBS-X available from Ciba Specialty Chemicals, 1,4-bis(2-cyanostyryl)benzene (C.I. Fluorescent Brightener 199) with tradename Ultraphor RN available from BASF.

Triazinstilbenes

Triazinstilbenes are a class of compounds having both triazin and stilbene structures in the same molecule.

Triazinstilbenes useful in the present invention include those having formulae (4):

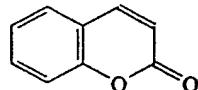


wherein R¹⁰⁷ and R¹⁰⁸, independently, are phenylamino, mono- or disulfonated phenylamino, morpholino, N(CH₂CH₂OH)₂, N(CH₃)(CH₂CH₂OH), NH₂, N(C₁-C₄-alkyl)₂, OCH₃, Cl, NH-(CH₂)₁₋₄SO₃H or NH-(CH₂)₁₋₄OH; An⁻ is an anion of a carboxylate, sulfate, sulfonate, or phosphate, and M is as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R¹⁰⁷ is 2, 5-disulfophenylamino and each R¹⁰⁸ is morpholino; or each R¹⁰⁷ is 2, 5-disulfophenylamino and each R¹⁰⁸ is N(C₂H₅)₂; or each R¹⁰⁷ is 3-sulfophenyl and each R¹⁰⁸ is NH(CH₂CH₂OH) or N(CH₂CH₂OH)₂; or each R¹⁰⁷ is 4-sulfophenyl and each R¹⁰⁸ is N(CH₂CH₂OH)₂; and in each case, the sulfo group is SO₃M in which M is sodium; wherein the compound has a trans-coplanar orientation.

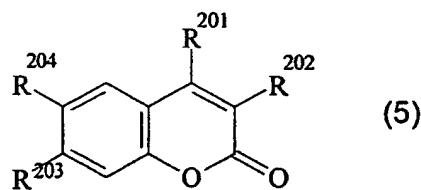
Suitable triazinstilbenes include 4,4'-bis-[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acid with tradename Tinopal UNPA-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4-anilino-6-morpholine-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate with tradename Tinopal AMS-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4-anilino-6-(2-hydroxyethyl)methyl amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate with tradename Tinopal 5BM-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4,6-dianilino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, 4,4'-bis-[(4-anilino-6-methylamino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, 4,4'-bis-[(4-anilino-6-ethylamino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, and 4,4'-bis(4-phenyl-1,2,3-triazol-2-yl)stilbene-2,2'-disulfonic acid.

Hydroxycoumarins

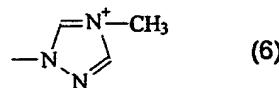
Hydroxycoumarins are a class of compounds having the following base coumarin structure and having at least one hydroxy moiety:



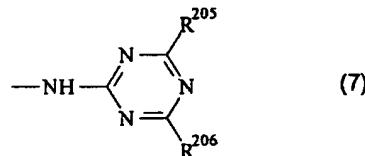
Hydroxycoumarins useful in the present invention include those having formulae (5):



wherein R²⁰¹ is H, OH, Cl, CH₃, CH₂COOH, CH₂SO₃H, CH₂OSO₃H, or CH₂OPO(OH)OH, R²⁰² is H, phenyl, COO-C₁-C₃₀-alkyl, glucose, or a group of formula (6):



and R²⁰³ is OH, or O-C₁-C₃₀-alkyl, and R²⁰⁴ is OH or O-C₁-C₃₀ alkyl, glycoside, or a group of the following formula (7):



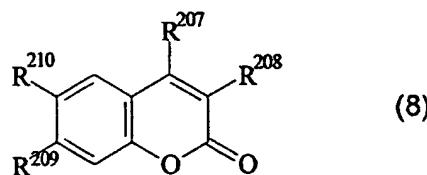
wherein R²⁰⁵ and R²⁰⁶ are independently, phenylamino, mono- or disulfonated phenylamino, morpholino, N(CH₂CH₂OH)₂, N(CH₃)(CH₂CH₂OH), NH₂, N(C₁-C₃₀-alkyl)₂, OCH₃, Cl, NH-(CH₂)₁₋₄SO₃H or NH-(CH₂)₁₋₄OH.

Suitable hydroxycoumarins include 6,7-dihydroxycoumarin available from Wako Chemicals, 4-methyl-7-hydroxycoumarin available from Wako Chemicals, 4-methyl-6,7-dihydroxycoumarin available from Wako Chemicals, esculin available from Wako Chemicals, and umbelliferone (4-hydroxycoumarin) available from Wako Chemicals.

Aminocoumarins

Aminocoumarins are a class of compounds having the base coumarin structure and having at least one amino moiety.

Aminocoumarins useful in the present inventions include those having formulae (8):

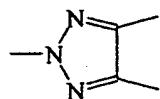


wherein R²⁰⁷ is H, Cl, CH₃, CH₂COOH, CH₂SO₃H, CH₂OSO₃H, or CH₂OPO(OH)OH, R²⁰⁸ is H, phenyl, or COOC₁-C₃₀ alkyl, and R²⁰⁹ and R²¹⁰ are independently H, NH₂, N(C₁-C₃₀alkyl)₂, NHC₁-C₃₀alkyl, or NHCOC₁-C₃₀alkyl.

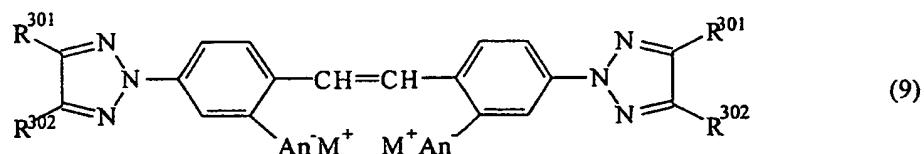
Suitable aminocoumarins include 4-methyl-7,7'-diethylamino coumarin with tradename Calcofluor-RWP available from BASF, 4-methyl-7,7'-dimethylamino coumarin with tradename Calcofluor-LD available from BASF.

Triazoles

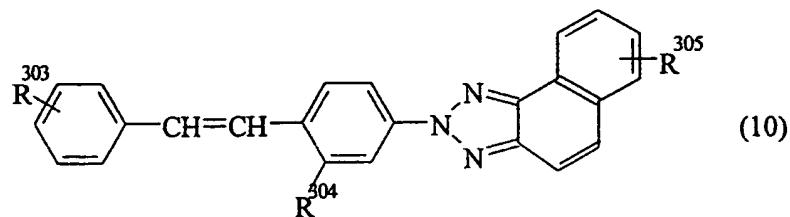
Triazoles are a class of compounds having the following base structure:



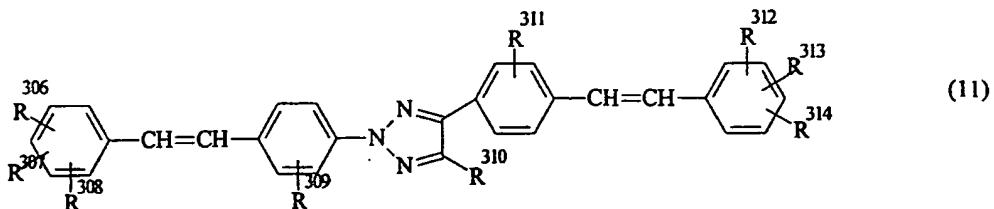
Triazoles useful in the present inventions include those having formulae (9) through (12) and (15) through (20):



wherein R³⁰¹ and R³⁰², independently, are H, C₁-C₃₀alkyl, phenyl or monosulfonated phenyl; An⁻ and M are as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R³⁰¹ is phenyl, R³⁰² is H and M is sodium; wherein the compound has a trans-coplanar orientation;

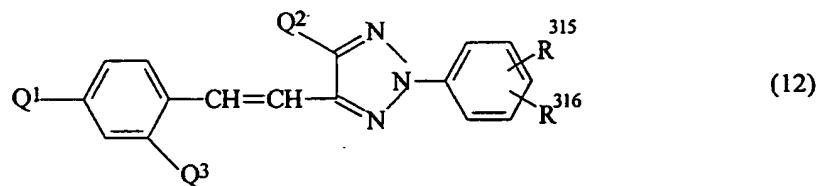


wherein R³⁰³ is H or Cl; R³⁰⁴ is SO₃M, SO₂N(C₁-C₃₀-alkyl)₂, SO₂O-phenyl or CN; R³⁰⁵ is H, SO₃M, COOM, OSO₃M, or OPO(OH)OM; and M is as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R³⁰³ and R³⁰⁵ are H and R³⁰⁴ is SO₃M in which M is Na; wherein the compound has a trans-coplanar orientation;

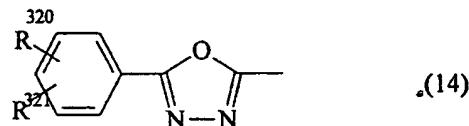
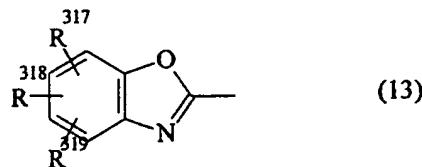


wherein each of R³⁰⁶ and R³¹² independently represents H, a sulfonic acid group or the salts, esters or amides thereof, a carboxylic acid group or the salts, esters or amides thereof, a cyano group, a halogen atom, an unsubstituted or substituted alkylsulfonyl, arylsulfonyl, alkyl, alkoxy, aralkyl, aryl, aryloxy, aralkoxy or cycloalkyl radical, an unsubstituted or substituted 5-membered heterocyclic ring containing 2 to 3 nitrogen atoms or one oxygen atom and 1 or 2 nitrogen atoms, or together with R³⁰⁷ and R³¹³ they represent a methylenedioxy, ethylenedioxy, methylenoxymethylenoxy, trimethylene, tetramethylene, propenylene, butenylene or butadienylene radical, each of R³⁰⁷ and R³¹³ independently represents H, a sulfonic acid group or the salts, esters or amides thereof, a carboxylic acid group or the salts, esters or amides thereof, a cyano group, a halogen atom, an unsubstituted or substituted alkyl or alkoxy radical, or together with R³⁰⁶ and R³¹² represent a methylenedioxy, ethylenedioxy, methylenoxymethylenoxy, trimethylene, tetramethylene, propenylene, butenylene or butadienylene radical, each of R³⁰⁸ and R³¹⁴ independently represents H, a halogen atom or an unsubstituted or substituted alkyl radical, each of R³⁰⁹ and R³¹¹ independently represents H, a halogen atom, a cyano group a sulonic acid group or the salts, esters or amides thereof, or a carboxylic acid group or the salts, esters or amides thereof, and R³¹⁰ independently represents H, a halogen atom, a cyano group a sulfonic acid group or the salts, alkyl radicals preferably by hydroxy, alkoxy of 1 to 30 carbon atoms, cyano, halogen, carboxy, sulfonic acid groups, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, phenyl or phenoxy; alkoxy radicals can be substituted by hydroxy, alkoxy of 1 to 30 carbon atoms, cyano, halogen, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, phenyl or phenoxy; phenyl, phenylalkyl or phenoxy radicals can be substituted by halogen, cyano, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, sulfo, or alkyl or alkoxy each of 1 to 30 carbon atoms; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; possible cycloalkyl radicals are preferably cyclohexyl and cyclopentyl radicals which can be substituted by alkyl

of 1 to 30 carbon atoms; possible 5-membered heterocyclic rings are v-triazole, oxazole or 1, 3, 4- oxadiazole radicals which can contain as substituents alkyl radicals of 1 to 4 carbon atoms, halogen, phenyl, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, cyano, benzyl, alkoxy of 1 to 30 carbon atoms, phenoxy or sulfo, whilst two adjacent substituents of the triazole and oxazole radicals together are able to form a substituted or unsubstituted fused benzene nucleus; wherein the compound has a trans-coplanar orientation;

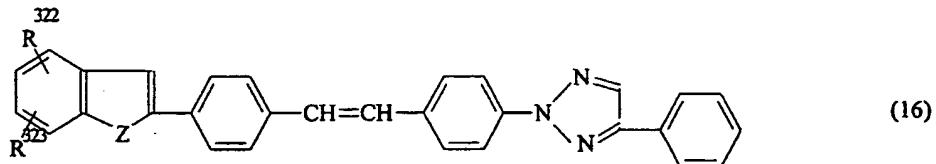
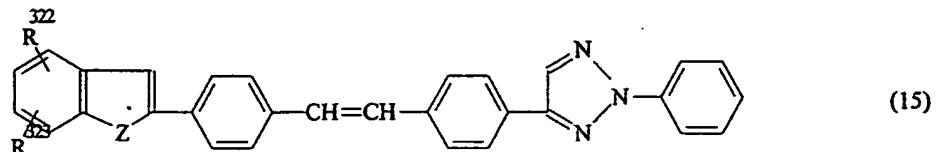


wherein Q^1 denotes one of the ring systems (13) or (14);

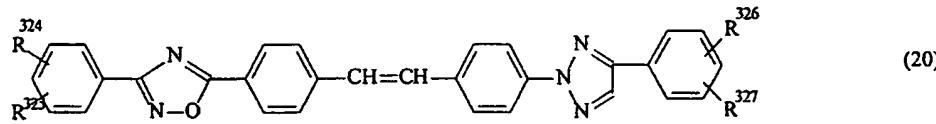
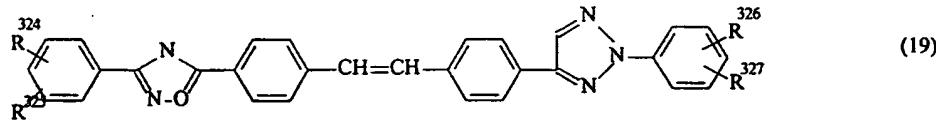
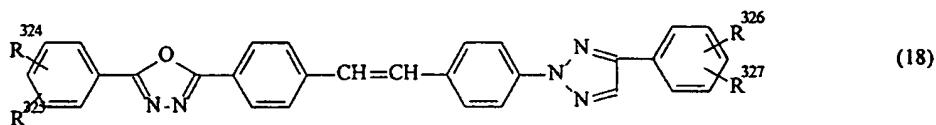
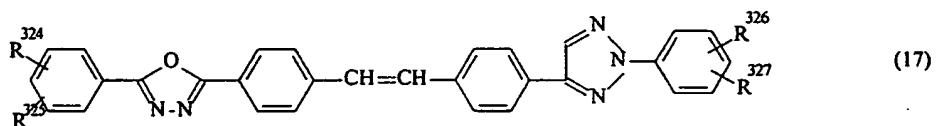


and wherein R^{317} denotes H, alkyl with 1 to 30 carbon atoms, cyclohexyl, phenylalkyl with C₁-C₃₀ carbon atoms in the alkyl part, phenyl, alkoxy with 1 to 30 carbon atoms, or Cl, or, conjointly with R^{318} , denotes alkylene with 3 to 30 carbon atoms, R^{318} denotes H or alkyl with 1 to 30 carbon atoms or, conjointly with R^{317} , denotes alkylene with 3 to 30 carbon atoms, R^{319} denotes H or methyl, R^{320} denotes H, alkyl with 1 to 30 carbon atoms, phenyl, alkoxy with 1 to 30 carbon atoms, or Cl, or, conjointly with R^{321} , denotes a fused benzene ring, R^{321} denotes H or Cl or conjointly with R^{320} , denotes a fused benzene ring, R^{315} denotes H, alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms or Cl, R^{316} denotes H or Cl, Q^2 denotes H, Cl alkyl with 1 to 30 carbon atoms or phenyl and Q^3 denotes H or Cl; wherein the compound has a

trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation;



wherein R322 denotes H, Cl, methyl, phenyl, benzyl, cyclohexyl or methoxy, R323 denotes H or methyl and Z denotes O or S; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation; and



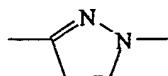
wherein R324 denotes H, Cl, alkyl with 1 to 30 carbon atoms, phenylalkyl with 1 to 30 carbon atoms, phenyl or alkoxy with 1 to 30 carbon atoms, or R324 conjointly with R325 denotes a fused benzene radical, R325 denotes H or methyl or R325 conjointly with R324 denotes a fused benzene radical, R326 denotes H, alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms, Cl, carbalkoxy with 1 to 30 carbon atoms or alkylsulfonyl with 1 to 30 carbon

atoms and R³²⁷ denotes H, Cl, methyl or methoxy; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

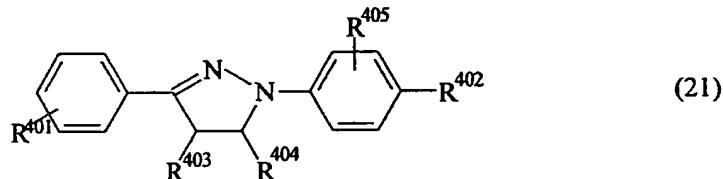
Suitable triazoles include 2-(4-styryl-3-sulfophenyl)-2H-naphtho[1,2-d]triazole (C.I. Fluorescent Brightener 46) with tradename Tinopal RBS available from Ciba Specialty Chemicals.

Pyrazolines

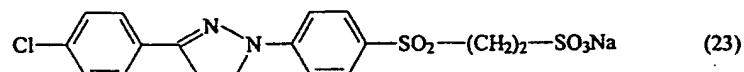
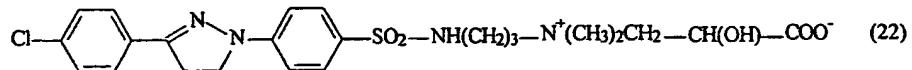
Pyrazolines are a class of compounds having the following base structure:



Pyrazolines useful in the present invention include those having formulae (21) through (23):



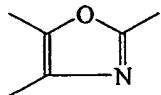
wherein R⁴⁰¹ is H, Cl or N(C₁-C₃₀-alkyl)₂, R⁴⁰² is H, Cl, SO₃M, SO₂NH₂, SO₂NH-(C₁-C₃₀alkyl), COO-C₁-C₃₀alkyl, SO₂-C₁-C₃₀alkyl, SO₂NH(CH₂)₁₋₄N⁺(CH₃)₃ or SO₂(CH₂)₁₋₄N⁺H(C₁-C₃₀-alkyl)₂An⁻, R⁴⁰³ and R⁴⁰⁴ are the same or different and each is H, C₁-C₃₀alkyl or phenyl and R⁴⁰⁵ is H or Cl; and An⁻ and M are as previously defined, preferably R⁴⁰¹ is Cl, R⁴⁰² is SO₂CH₂CH₂N⁺H(C₁-C₄-alkyl)₂An⁻ in which An⁻ is phosphite and R⁴⁰³, R⁴⁰⁴ and R⁴⁰⁵ are each H; and formulae (22) and (23) shown below.



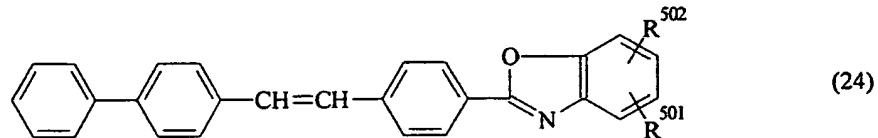
Suitable pyrazolines include 1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline (C.I. Fluorescent Brightener 121) with tradename Blankophor DCB available from Bayer, 1-[4-(2-sulfoethylsulfonyl)phenyl]-3-(4-chlorophenyl)-2-pyrazoline, 1-[4-(2-sulfoethylsulfonyl)phenyl]-3-(3,4-dichloro-6-methylphenyl)-2-pyrazoline, 1-<4-{N-[3-(N,N,N-trimethylammonio)propyl]-amidosulfonyl}phenyl>-3-(4-chlorophenyl)-2-pyrazoline methylsulfate, and 1-<4-{2-[1-methyl-2-(N,N-dimethylamino)ethoxy]ethylsulfonyl}phenyl>-3-(4-chlorophenyl)-2-pyrazoline methylsulfate.

Oxazoles

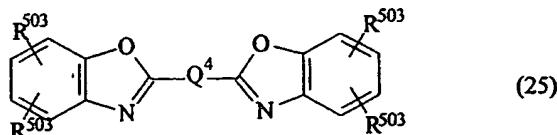
Oxazoles are a class of compounds having the following base structure:



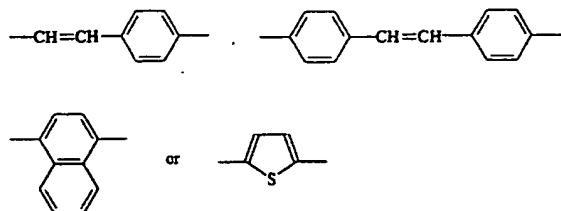
Oxazoles useful in the present inventions include those having formulae (24), (25), (26) and (27):



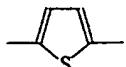
wherein R⁵⁰¹ and R⁵⁰², independently, are H, Cl, C₁-C₃₀alkyl or SO₂-C₁-C₃₀alkyl, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R⁵⁰¹ is 4-CH₃ and R⁵⁰² is 2-CH₃ wherein the compound has a trans-coplanar orientation;



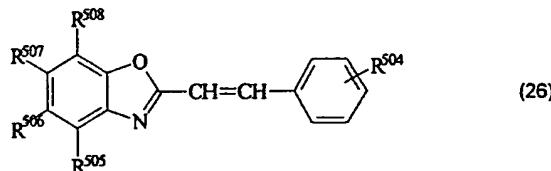
wherein R⁵⁰³, independently, is H, C(CH₃)₃, C(CH₃)₂-phenyl, C₁-C₃₀alkyl or COO-C₁-C₃₀alkyl, preferably H and Q⁴ is -CH=CH-;



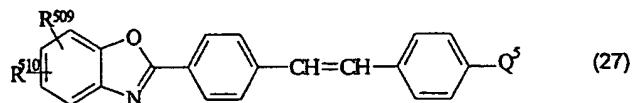
preferably



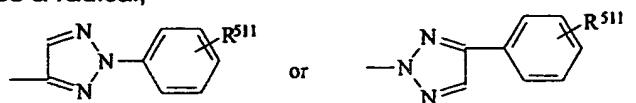
or one group R⁵⁰³ in each ring is 2-methyl and the other R⁵⁰³ is H and Q⁴ is -CH=CH-; or one group R⁵⁰³ in each ring is 2-C(CH₃)₃ and the other R⁵⁰³ is H; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation;



wherein R⁵⁰⁴ is CN, Cl, COO-C₁-C₃₀alkyl or phenyl; R⁵⁰⁵ and R⁵⁰⁸ are the atoms required to form a fused benzene ring or R⁵⁰⁶ and R⁵⁰⁸, independently, are H or C₁-C₃₀alkyl; and R⁵⁰⁷ is H, C₁-C₃₀alkyl or phenyl; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R⁵⁰⁴ is a 4-phenyl group and each of R⁵⁰⁵ to R⁵⁰⁸ is H; wherein the compound has a trans-coplanar orientation; and



wherein R⁵⁰⁹ denotes H, Cl, alkyl with 1 to 30 carbon atoms, cyclohexyl, phenylalkyl with 1 to 3 carbon atoms in the alkyl part, phenyl or alkoxy with 1 to 30 carbon atoms, R⁵¹⁰ denotes H or alkyl with 1 to 30 carbon atoms, and Q⁵ denotes a radical;

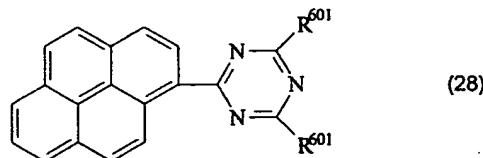


wherein R⁵¹¹ represents H, alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms, Cl, carbalkoxy with 1 to 30 carbon atoms, unsubstituted sulfamoyl or sulfamoyl which is monosubstituted or disubstituted by alkyl or hydroxyalkyl with 1 to 30 carbon atoms or represents alkylsulfonyl with 1 to 30 carbon atoms; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

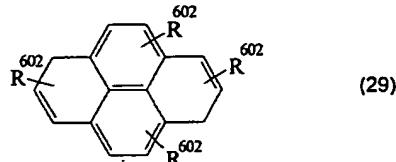
Suitable oxazoles include 4,4'-bis(5-methylbenzoxazol-2-yl)stilbene, and 2-(4-methoxycarbonylstyryl)benzoxazole.

Pyrenes

Pyrenes useful in the present invention include those having formulae (28) and (29):



wherein each R⁶⁰¹, independently, is C1-C30alkoxy; preferably methoxy; and

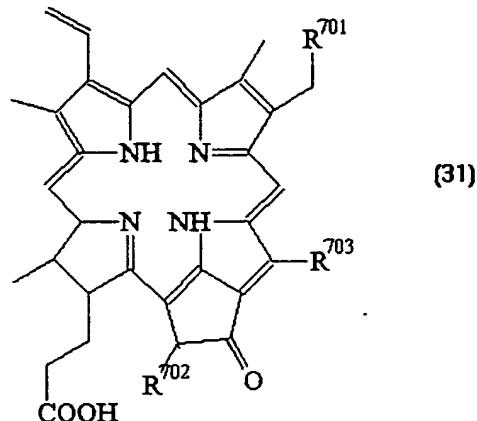
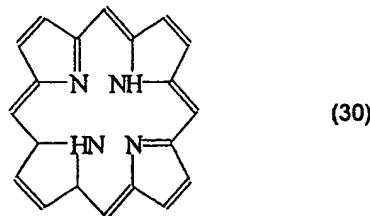


wherein each R⁶⁰², independently, is H, OH, or SO₃M, wherein M is as previously defined, sulfonated phenylamino, or anilino.

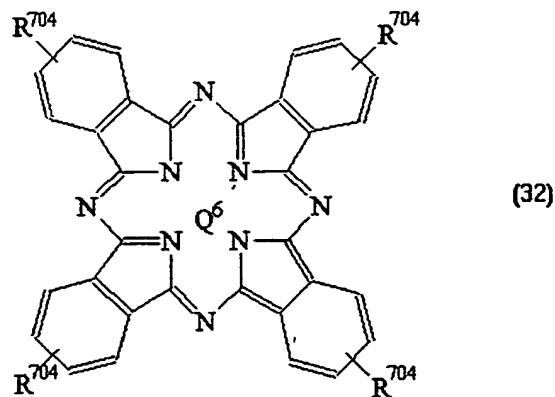
Suitable pyrenes include 2,4-dimethoxy-6-(1'-pyrenyl)-1,3,5-triazine (C.I. Fluorescent Brightener 179) with tradename Fluolite XMF, 8-hydroxy-1,3,6-pyrenetrisulfonic acid (D&C Green No.8), and 3-hydroxy-5,8,10-trisulphanilic pyrene.

Porphyrins

Porphyrins useful in the present invention include those having formulae (30), (31), and (32):



wherein R701 is CH₃ or CHO, R702 is H or COOC₁-C₃₀ alkyl, and R703 is H or an alkyl group having 1 to 30 carbons; and

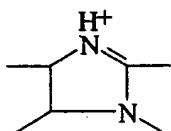


wherein each R704, independently, is H, SO₃M, COOM, OSO₃M, or OPO(OH)OM, wherein M is as previously defined, halide, or alkyl of 1 to 30 carbons; and Q6 is Cu, Mg, Fe, Cr, Co, or mixtures thereof with cationic charges.

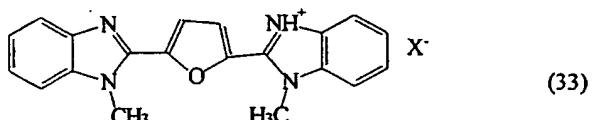
Suitable porphyrins include porphyrin available from Wako Chemicals and Copper II phthalocyanine available from Wako Chemicals.

Imidazoles

Imidazoles are a class of compounds having the following base structure:



Imidazoles useful in the present invention include those having formulae (33):



wherein X- is as previously defined.

Suitable imidazoles include those with tradename of C.I. Fluorescence Brightener 352, or Uvtex AT available from Ciba Speciality Chemical.

COMPOSITIONS

The hair care compositions of the present invention may comprise a component which characterizes the form of the product. Product forms useful herein include, but are not limited to, shampoo, conditioner, treatment, mousse, spray, lotion, gel, and cream products, all of which can be designed for rinse-off or leave-on convenience. Product functions useful herein include, but are not limited to, cleansing, and conditioning products. The components and their levels are selected by one skilled in the art depending on the desired characteristic of the product.

HAIR CONDITIONING AGENT

The compositions of the present invention comprise a hair conditioning agent selected from the group consisting of silicone compounds, cationic compounds, water-insoluble high melting point compounds, perfume compounds, high molecular weight oily compounds, and mixtures thereof. The hair conditioning compositions herein provide good conditioning benefits such as good combing, smoothness, and reduction of static. Preferably, the hair

conditioning agent is comprised by weight in the present composition at a level of from about 0.01% to about 20%.

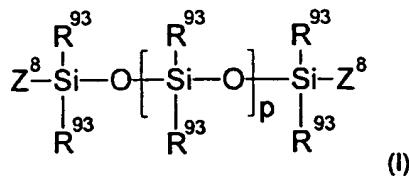
Silicone Compound

Silicone compounds are useful hair conditioning agents herein, and are preferably include at a level of from about 0.01% to about 15%. The silicone compounds useful herein include volatile soluble or insoluble, or nonvolatile soluble or insoluble silicone conditioning agents. By soluble what is meant is that the silicone compound is miscible with the carrier of the composition so as to form part of the same phase. By insoluble what is meant is that the silicone forms a separate, discontinuous phase from the carrier, such as in the form of an emulsion or a suspension of droplets of the silicone. The silicone compounds herein may be made by any suitable method known in the art, including emulsion polymerization. The silicone compounds may further be incorporated in the present composition in the form of an emulsion, wherein the emulsion is made my mechanical mixing, or in the stage of synthesis through emulsion polymerization, with or without the aid of a surfactant selected from anionic surfactants, nonionic surfactants, cationic surfactants, and mixtures thereof.

In one embodiment, the silicone compound useful herein is a high molecular weight silicone compound. The high molecular weight silicone compounds useful herein have a viscosity of from about 1,000 to about 2,000,000 centistokes at 25°C, more preferably from about 10,000 to about 1,800,000, and even more preferably from about 100,000 to about 1,500,000. The viscosity can be measured by means of a glass capillary viscometer as set forth in Dow Corning Corporate Test Method CTM0004, July 20, 1970. High molecular weight silicone compounds may be made by emulsion polymerization. Suitable silicone fluids include polyalkyl siloxanes, polyaryl siloxanes, polyalkylaryl siloxanes, polyether siloxane copolymers, and mixtures thereof. Other nonvolatile silicone compounds having hair conditioning properties can also be used.

The silicone compounds herein are preferably used at levels by weight of the composition of from about 0.1% to about 60%, more preferably from about 0.1% to about 40%.

The silicone compounds herein also include polyalkyl or polyaryl siloxanes with the following structure (I)



wherein R^{93} is alkyl or aryl, and x is an integer from about 7 to about 8,000. Z^8 represents groups which block the ends of the silicone chains. The alkyl or aryl groups substituted on the siloxane chain (R^{93}) or at the ends of the siloxane chains Z^8 can have any structure as long as the resulting silicone remains fluid at room temperature, is dispersible, is neither irritating, toxic nor otherwise harmful when applied to the hair, is compatible with the other components of the composition, is chemically stable under normal use and storage conditions, and is capable of being deposited on and conditions the hair. Suitable Z^8 groups include hydroxy, methyl, methoxy, ethoxy, propoxy, and aryloxy. The two R^{93} groups on the silicon atom may represent the same group or different groups. Preferably, the two R^{93} groups represent the same group. Suitable R^{93} groups include methyl, ethyl, propyl, phenyl, methylphenyl and phenylmethyl. The preferred silicone compounds are polydimethylsiloxane, polydiethylsiloxane, and polymethylphenylsiloxane. Polydimethylsiloxane, which is also known as dimethicone, is especially preferred. The polyalkylsiloxanes that can be used include, for example, polydimethylsiloxanes. These silicone compounds are available, for example, from the General Electric Company in their Viscasil® and SF 96 series, and from Dow Corning in their Dow Corning 200 series.

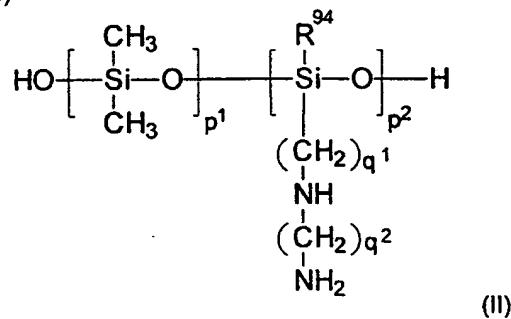
Polyalkylaryl siloxane fluids can also be used and include, for example, polymethylphenylsiloxanes. These siloxanes are available, for example, from the General Electric Company as SF 1075 methyl phenyl fluid or from Dow Corning as 556 Cosmetic Grade Fluid.

Especially preferred, for enhancing the shine characteristics of hair, are highly arylated silicone compounds, such as highly phenylated polyethyl silicone having refractive index of about 1.46 or higher, especially about 1.52 or higher. When these high refractive index silicone compounds are used, they should be mixed with a spreading agent, such as a surfactant or a silicone resin, as described below to decrease the surface tension and enhance the film forming ability of the material.

The silicone compounds that can be used include, for example, a polypropylene oxide modified polydimethylsiloxane although ethylene oxide or mixtures of ethylene oxide and propylene oxide can also be used. The ethylene

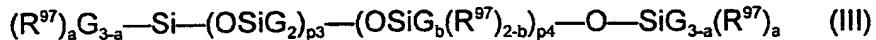
oxide and polypropylene oxide level should be sufficiently low so as not to interfere with the dispersibility characteristics of the silicone. These material are also known as dimethicone copolyols.

Other silicone compounds include amino substituted materials. Suitable alkylamino substituted silicone compounds include those represented by the following structure (II)

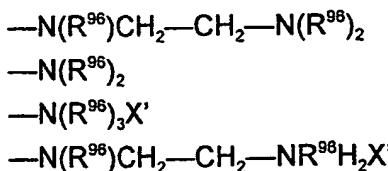


wherein R^{94} is H, CH_3 or OH , p^1 , p^2 , q^1 and q^2 are integers which depend on the molecular weight, the average molecular weight being approximately between 5,000 and 10,000. This polymer is also known as "amodimethicone".

Suitable amino substituted silicone fluids include those represented by the formula (III)



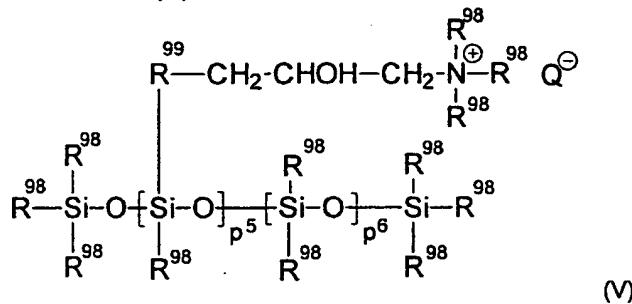
in which G is chosen from the group consisting of hydrogen, phenyl, OH, $\text{C}_1\text{-C}_8$ alkyl and preferably methyl; a denotes 0 or an integer from 1 to 3, and preferably equals 0; b denotes 0 or 1 and preferably equals 1; the sum p^3+p^4 is a number from 1 to 2,000 and preferably from 50 to 150, p^3 being able to denote a number from 0 to 1,999 and preferably from 49 to 149 and p^4 being able to denote an integer from 1 to 2,000 and preferably from 1 to 10; R^{97} is a monovalent radical of formula $\text{C}_{q^3}\text{H}_{2q^3}\text{L}$ in which q^3 is an integer from 2 to 8 and L is chosen from the groups



in which R^{98} is chosen from the group consisting of hydrogen, phenyl, benzyl, a saturated hydrocarbon radical, preferably an alkyl radical containing from 1 to 20 carbon atoms, and X' denotes a halide ion.

An especially preferred amino substituted silicone corresponding to formula (II) is the polymer known as "trimethylsilylamodimethicone" wherein R⁹⁴ is CH₃.

Other amino substituted silicone polymers which can be used are represented by the formula (V):



where R⁹⁸ denotes a monovalent hydrocarbon radical having from 1 to 18 carbon atoms, preferably an alkyl or alkenyl radical such as methyl; R⁹⁹ denotes a hydrocarbon radical, preferably a C₁-C₁₈ alkylene radical or a C₁-C₁₈, and more preferably C₁-C₈, alkyleneoxy radical; Q[⊖] is a halide ion, preferably chloride; p⁵ denotes an average statistical value from 2 to 20, preferably from 2 to 8; p⁶ denotes an average statistical value from 20 to 200, and preferably from 20 to 50. A preferred polymer of this class is available from Union Carbide under the name "UCAR SILICONE ALE 56."

References disclosing suitable nonvolatile dispersed silicone compounds include U.S. Patent No. 2,826,551, to Geen; U.S. Patent No. 3,964,500, to Drakoff, issued June 22, 1976; U.S. Patent No. 4,364,837, to Pader; and British Patent No. 849,433, to Woolston. "Silicon Compounds" distributed by Petrarch Systems, Inc., 1984, provides an extensive, though not exclusive, listing of suitable silicone compounds.

Another nonvolatile dispersed silicone that can be especially useful is a silicone gum. The term "silicone gum", as used herein, means a polyorganosiloxane material having a viscosity at 25°C of greater than or equal to 1,000,000 centistokes. It is recognized that the silicone gums described herein can also have some overlap with the above-disclosed silicone compounds. This overlap is not intended as a limitation on any of these materials. Silicone gums are described by Petrarch, and others including U.S. Patent No. 4,152,416, to Spitzer *et al.*, issued May 1, 1979 and Noll, Walter, Chemistry and Technology of Silicones, New York: Academic Press 1968. Also describing silicone gums are General Electric Silicone Rubber Product Data

Sheets SE 30, SE 33, SE 54 and SE 76. The "silicone gums" will typically have a mass molecular weight in excess of about 200,000, generally between about 200,000 and about 1,000,000. Specific examples include polydimethylsiloxane, poly(dimethylsiloxane methylvinylsiloxane) copolymer, poly(dimethylsiloxane diphenylsiloxane methylvinylsiloxane) copolymer and mixtures thereof.

Also useful are silicone resins, which are highly crosslinked polymeric siloxane systems. The crosslinking is introduced through the incorporation of tri-functional and tetra-functional silanes with mono-functional or di-functional, or both, silanes during manufacture of the silicone resin. As is well understood in the art, the degree of crosslinking that is required in order to result in a silicone resin will vary according to the specific silane units incorporated into the silicone resin. In general, silicone materials which have a sufficient level of trifunctional and tetrafunctional siloxane monomer units, and hence, a sufficient level of crosslinking, such that they dry down to a rigid, or hard, film are considered to be silicone resins. The ratio of oxygen atoms to silicon atoms is indicative of the level of crosslinking in a particular silicone material. Silicone materials which have at least about 1.1 oxygen atoms per silicon atom will generally be silicone resins herein. Preferably, the ratio of oxygen:silicon atoms is at least about 1.2:1.0. Silanes used in the manufacture of silicone resins include monomethyl-, dimethyl-, trimethyl-, monophenyl-, diphenyl-, methylphenyl-, monovinyl-, and methylvinylchlorosilanes, and tetrachlorosilane, with the methyl substituted silanes being most commonly utilized. Preferred resins are offered by General Electric as GE SS4230 and SS4267. Commercially available silicone resins will generally be supplied in a dissolved form in a low viscosity volatile or nonvolatile silicone fluid. The silicone resins for use herein should be supplied and incorporated into the present compositions in such dissolved form, as will be readily apparent to those skilled in the art. Without being bound by theory, it is believed that the silicone resins can enhance deposition of other silicone compounds on the hair and can enhance the glossiness of hair with high refractive index volumes.

Other useful silicone resins are silicone resin powders such as the material given the CTFA designation polymethylsilsequioxane, which is commercially available as Tospearl™ from Toshiba Silicones.

The method of manufacturing these silicone compounds, can be found in Encyclopedia of Polymer Science and Engineering, Volume 15, Second Edition, pp. 204-308, John Wiley & Sons, Inc., 1989.

Silicone materials and silicone resins in particular, can conveniently be identified according to a shorthand nomenclature system well known to those skilled in the art as the "MDTQ" nomenclature. Under this system, the silicone is described according to the presence of various siloxane monomer units which make up the silicone. Briefly, the symbol M denotes the mono-functional unit $(CH_3)_3SiO_{0.5}$; D denotes the difunctional unit $(CH_3)_2SiO$; T denotes the trifunctional unit $(CH_3)SiO_{1.5}$; and Q denotes the quadri- or tetra-functional unit SiO_2 . Primes of the unit symbols, e.g., M', D', T', and Q' denote substituents other than methyl, and must be specifically defined for each occurrence. Typical alternate substituents include groups such as vinyl, phenyl, amino, hydroxyl, etc. The molar ratios of the various units, either in terms of subscripts to the symbols indicating the total number of each type of unit in the silicone, or an average thereof, or as specifically indicated ratios in combination with molecular weight, complete the description of the silicone material under the MDTQ system. Higher relative molar amounts of T, Q, T' and/or Q' to D, D', M and/or or M' in a silicone resin is indicative of higher levels of crosslinking. As discussed before, however, the overall level of crosslinking can also be indicated by the oxygen to silicon ratio.

The silicone resins for use herein which are preferred are MQ, MT, MTQ, MQ and MDTQ resins. Thus, the preferred silicone substituent is methyl. Especially preferred are MQ resins wherein the M:Q ratio is from about 0.5:1.0 to about 1.5:1.0 and the average molecular weight of the resin is from about 1000 to about 10,000.

Particularly suitable silicone compounds herein are non-volatile silicone oils having a molecular weight of from about 200,000 to about 600,000 such as Dimethicone, and Dimethiconol. These silicone compounds can be incorporated in the composition as silicone oils solutions; the silicone oils being volatile or non-volatile.

Commercially available silicone compounds which are useful herein include Dimethicone with tradename DC345 available from Dow Corning Corporation, Dimethicone gum solutions with tradenames SE 30, SE 33, SE 54 and SE 76 available from General Electric, Dimethiconol with tradenames DCQ2-1403 and DCQ2-1401 available from Dow Corning Corporation, and emulsion polymerized Dimethiconol available from Toshiba Silicone as described in GB application 2,303,857.

Cationic Compounds

The cationic compounds useful herein include cationic polymers and cationic surfactants. Preferably, the cationic compound has no fluorescent property, and is present at a level of from about 0.01% to about 10%.

Cationic Polymers

As used herein, the term "polymer" shall include materials whether made by polymerization of one type of monomer or made by two (i.e., copolymers) or more types of monomers.

Preferably, the cationic polymer is a water-soluble cationic polymer. By "water soluble" cationic polymer, what is meant is a polymer which is sufficiently soluble in water to form a substantially clear solution to the naked eye at a concentration of 0.1% in water (distilled or equivalent) at 25°C. The preferred polymer will be sufficiently soluble to form a substantially clear solution at 0.5% concentration, more preferably at 1.0% concentration.

The cationic polymers hereof will generally have a weight average molecular weight which is at least about 5,000, typically at least about 10,000, and is less than about 10 million. Preferably, the molecular weight is from about 100,000 to about 2 million. The cationic polymers will generally have cationic nitrogen-containing moieties such as quaternary ammonium or cationic amino moieties, and mixtures thereof.

The cationic charge density is preferably at least about 0.1 meq/gram, more preferably at least about 1.5 meq/gram, even more preferably at least about 1.1 meq/gram, still more preferably at least about 1.2 meq/gram. Cationic charge density of the cationic polymer can be determined according to the Kjeldahl Method. Those skilled in the art will recognize that the charge density of amino-containing polymers may vary depending upon pH and the isoelectric point of the amino groups. The charge density should be within the above limits at the pH of intended use.

Any anionic counterions can be utilized for the cationic polymers so long as the water solubility criteria is met. Suitable counterions include halides (e.g., Cl, Br, I, or F, preferably Cl, Br, or I), sulfate, and methylsulfate. Others can also be used, as this list is not exclusive.

The cationic nitrogen-containing moiety will be present generally as a substituent, on a fraction of the total monomer units of the cationic hair conditioning polymers. Thus, the cationic polymer can comprise copolymers, terpolymers, etc. of quaternary ammonium or cationic amine-substituted

monomer units and other non-cationic units referred to herein as spacer monomer units. Such polymers are known in the art, and a variety can be found in the CTFA Cosmetic Ingredient Dictionary, 3rd edition, edited by Estrin, Crosley, and Haynes, (The Cosmetic, Toiletry, and Fragrance Association, Inc., Washington, D.C., 1982).

Suitable cationic polymers include, for example, copolymers of vinyl monomers having cationic amine or quaternary ammonium functionalities with water soluble spacer monomers such as acrylamide, methacrylamide, alkyl and dialkyl acrylamides, alkyl and dialkyl methacrylamides, alkyl acrylate, alkyl methacrylate, vinyl caprolactone, and vinyl pyrrolidone. The alkyl and dialkyl substituted monomers preferably have C₁ - C₇ alkyl groups, more preferably C₁ - C₃ alkyl groups. Other suitable spacer monomers include vinyl esters, vinyl alcohol (made by hydrolysis of polyvinyl acetate), maleic anhydride, propylene glycol, and ethylene glycol.

The cationic amines can be primary, secondary, or tertiary amines, depending upon the particular species and the pH of the composition. In general, secondary and tertiary amines, especially tertiary amines, are preferred.

Amine-substituted vinyl monomers can be polymerized in the amine form, and then optionally can be converted to ammonium by a quaternization reaction. Amines can also be similarly quaternized subsequent to formation of the polymer. For example, tertiary amine functionalities can be quaternized by reaction with a salt of the formula R'X wherein R' is a short chain alkyl, preferably a C₁ - C₇ alkyl, more preferably a C₁ - C₃ alkyl, and X is an anion which forms a water soluble salt with the quaternized ammonium.

Suitable cationic amino and quaternary ammonium monomers include, for example, vinyl compounds substituted with dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, monoalkylaminoalkyl acrylate, monoalkylaminoalkyl methacrylate, trialkyl methacryloxyalkyl ammonium salt, trialkyl acryloxyalkyl ammonium salt, diallyl quaternary ammonium salts, and vinyl quaternary ammonium monomers having cyclic cationic nitrogen-containing rings such as pyridinium, imidazolium, and quaternized pyrrolidone, e.g., alkyl vinyl imidazolium, alkyl vinyl pyridinium, alkyl vinyl pyrrolidone salts. The alkyl portions of these monomers are preferably lower alkyls such as the C₁ - C₃ alkyls, more preferably C₁ and C₂ alkyls. Suitable amine-substituted vinyl monomers for use herein include dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, dialkylaminoalkyl acrylamide, and dialkylaminoalkyl

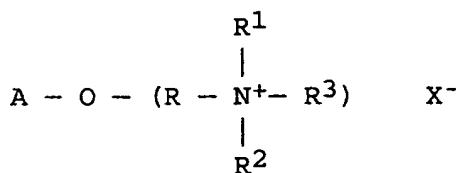
methacrylamide, wherein the alkyl groups are preferably C₁ - C₇ hydrocarbyls, more preferably C₁ - C₃, alkyls.

The cationic polymers hereof can comprise mixtures of monomer units derived from amine- and/or quaternary ammonium-substituted monomer and/or compatible spacer monomers.

Suitable cationic hair conditioning polymers include, for example: copolymers of 1-vinyl-2-pyrrolidone and 1-vinyl-3-methylimidazolium salt (e.g., chloride salt) (referred to in the industry by the Cosmetic, Toiletry, and Fragrance Association, "CTFA", as Polyquaternium-16), such as those commercially available from BASF Wyandotte Corp. (Parsippany, NJ, USA) under the LUVIQUAT tradename (e.g., LUVIQUAT FC 370); copolymers of 1-vinyl-2-pyrrolidone and dimethylaminoethyl methacrylate (referred to in the industry by CTFA as Polyquaternium-11) such as those commercially available from Gaf Corporation (Wayne, NJ, USA) under the GAFQUAT tradename (e.g., GAFQUAT 755N); cationic diallyl quaternary ammonium-containing polymers, including, for example, dimethyldiallylammonium chloride homopolymer and copolymers of acrylamide and dimethyldiallylammonium chloride, referred to in the industry (CTFA) as Polyquaternium 6 and Polyquaternium 7, respectively; and mineral acid salts of amino-alkyl esters of homo- and co-polymers of unsaturated carboxylic acids having from 3 to 5 carbon atoms, as described in U.S. Patent 4,009,256, incorporated herein by reference.

Other cationic polymers that can be used include polysaccharide polymers, such as cationic cellulose derivatives and cationic starch derivatives.

Cationic polysaccharide polymer materials suitable for use herein include those of the formula:



wherein: A is an anhydroglucose residual group, such as a starch or cellulose anhydroglucose residual, R is an alkylene oxyalkylene, polyoxyalkylene, or hydroxyalkylene group, or combination thereof, R¹, R², and R³ independently are alkyl, aryl, alkylaryl, arylalkyl, alkoxyalkyl, or alkoxyaryl groups, each group containing up to about 18 carbon atoms, and the total number of carbon atoms for each cationic moiety (i.e., the sum of carbon atoms in R¹, R² and R³)

preferably being about 20 or less, and X is an anionic counterion, as previously described.

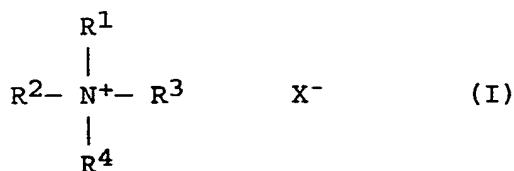
Cationic cellulose is available from Amerchol Corp. (Edison, NJ, USA) in their Polymer JR® and LR® series of polymers, as salts of hydroxyethyl cellulose reacted with trimethyl ammonium substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 10. Another type of cationic cellulose includes the polymeric quaternary ammonium salts of hydroxyethyl cellulose reacted with lauryl dimethyl ammonium-substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 24. These materials are available from Amerchol Corp. (Edison, NJ, USA) under the tradename Polymer LM-200®.

Other cationic polymers that can be used include cationic guar gum derivatives, such as guar hydroxypropyltrimonium chloride (commercially available from Celanese Corp. in their Jaguar R series). Other materials include quaternary nitrogen-containing cellulose ethers (e.g., as described in U.S. Patent 3,962,418, incorporated herein by reference), and copolymers of etherified cellulose and starch (e.g., as described in U.S. Patent 3,958,581, incorporated herein by reference.)

Cationic Surfactant

The cationic surfactants useful herein are any known to the artisan.

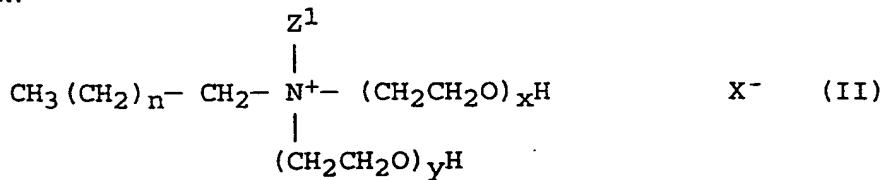
Among the cationic surfactants useful herein are those corresponding to the general formula (I):



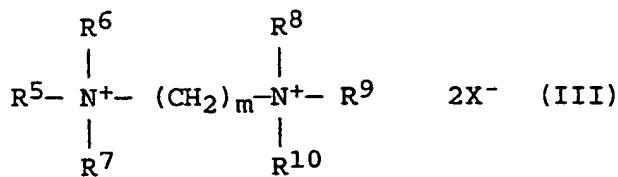
wherein at least one of R¹, R², R³, and R⁴ is selected from an aliphatic group of from 8 to 30 carbon atoms or an aromatic, alkoxy, polyoxyalkylene, alkylamido, hydroxyalkyl, aryl or alkylaryl group having up to about 22 carbon atoms, the remainder of R¹, R², R³, and R⁴ are independently selected from an aliphatic group of from 1 to about 22 carbon atoms or an aromatic, alkoxy, polyoxyalkylene, alkylamido, hydroxyalkyl, aryl or alkylaryl group having up to about 22 carbon atoms; and X is a salt-forming anion such as those selected from halogen, (e.g. chloride, bromide), acetate, citrate, lactate, glycolate, phosphate, nitrate, sulfonate, sulfate, alkylsulfate, and alkyl sulfonate radicals.

The aliphatic groups can contain, in addition to carbon and hydrogen atoms, ether linkages, and other groups such as amino groups. The longer chain aliphatic groups, e.g., those of about 12 carbons, or higher, can be saturated or unsaturated. Preferred is when R¹, R², R³, and R⁴ are independently selected from C₁ to about C₂₂ alkyl. Nonlimiting examples of cationic surfactants useful include the materials having the following CTFA designations: quaternium-8, quaternium-24, quaternium-26, quaternium-27, quaternium-30, quaternium-33, quaternium-43, quaternium-52, quaternium-53, quaternium-56, quaternium-60, quaternium-62, quaternium-70, quaternium-72, quaternium-75, quaternium-77, quaternium-78, quaternium-80, quaternium-81, quaternium-82, quaternium-83, quaternium-84, and mixtures thereof.

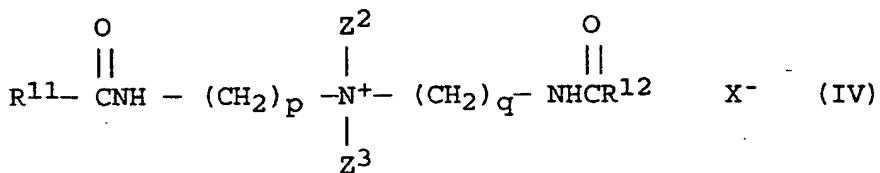
Also preferred are hydrophilically substituted cationic surfactants in which at least one of the substituents contain one or more aromatic, ether, ester, amido, or amino moieties present as substituents or as linkages in the radical chain, wherein at least one of the R¹ - R⁴ radicals contain one or more hydrophilic moieties selected from alkoxy (preferably C₁ - C₃ alkoxy), polyoxyalkylene (preferably C₁ - C₃ polyoxyalkylene), alkylamido, hydroxyalkyl, alkylester, and combinations thereof. Preferably, the hydrophilically substituted cationic conditioning surfactant contains from 2 to about 10 nonionic hydrophile moieties located within the above stated ranges. Preferred hydrophilically substituted cationic surfactants include those of the formula (II) through (VII) below:



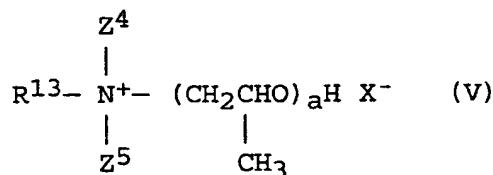
wherein n is from 8 to about 28, x+y is from 2 to about 40, Z¹ is a short chain alkyl, preferably a C₁ - C₃ alkyl, more preferably methyl, or -(CH₂CH₂O)_zH wherein x+y+z is up to 60, and X is a salt forming anion as defined above;



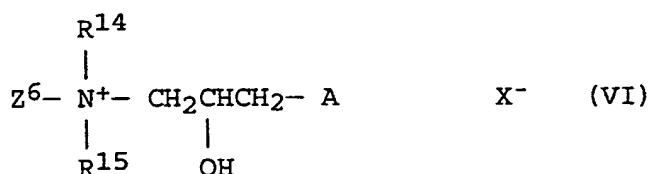
wherein m is 1 to 5, one or more of R⁵, R⁶, and R⁷ are independently an C₁ - C₃₀ alkyl, the remainder are -CH₂CH₂OH, one or two of R⁸, R⁹, and R¹⁰ are independently an C₁ - C₃₀ alkyl, and remainder are -CH₂CH₂OH, and X is a salt forming anion as mentioned above;



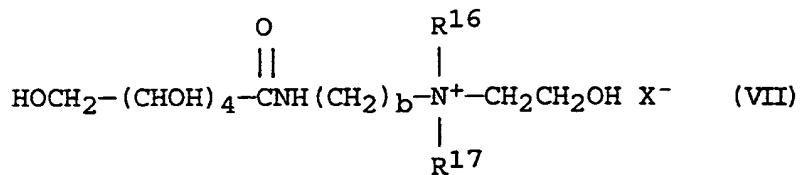
wherein Z² is an alkyl, preferably a C₁ - C₃ alkyl, more preferably methyl, and Z³ is a short chain hydroxyalkyl, preferably hydroxymethyl or hydroxyethyl, p and q independently are integers from 2 to 4, inclusive, preferably from 2 to 3, inclusive, more preferably 2, R¹¹ and R¹², independently, are substituted or unsubstituted hydrocarbyls, preferably C₁₂ - C₂₀ alkyl or alkenyl, and X is a salt forming anion as defined above;



wherein R¹³ is a hydrocarbyl, preferably a C₁ - C₃ alkyl, more preferably methyl, Z⁴ and Z⁵ are, independently, short chain hydrocarbyls, preferably C₂ - C₄ alkyl or alkenyl, more preferably ethyl, a is from 2 to about 40, preferably from about 7 to about 30, and X is a salt forming anion as defined above;



wherein R¹⁴ and R¹⁵, independently, are C₁ - C₃ alkyl, preferably methyl, Z⁶ is a C₁₂ - C₂₂ hydrocarbyl, alkyl carboxy or alkylamido, and A is a protein, preferably a collagen, keratin, milk protein, silk, soy protein, wheat protein, or hydrolyzed forms thereof, and X is a salt forming anion as defined above;



wherein b is 2 or 3, R¹⁶ and R¹⁷, independently are C₁ - C₃ hydrocarbys preferably methyl, and X is a salt forming anion as defined above. Nonlimiting examples of hydrophilically substituted cationic surfactants useful include the materials having the following CTFA designations: quaternium-16, quaternium-61, quaternium-71, quaternium-79 hydrolyzed collagen, quaternium-79 hydrolyzed keratin, quaternium-79 hydrolyzed milk protein, quaternium-79 hydrolyzed silk, quaternium-79 hydrolyzed soy protein, and quaternium-79 hydrolyzed wheat protein. Highly preferred compounds include commercially available materials of the following tradenames; VARIQUAT K1215 and 638 from Witco Chemical, MACKPRO KLP, MACKPRO WLW, MACKPRO MLP, MACKPRO NSP, MACKPRO NLW, MACKPRO WWP, MACKPRO NLP, MACKPRO SLP from McIntyre, ETHOQUAD 18/25, ETHOQUAD O/12PG, ETHOQUAD C/25, ETHOQUAD S/25, and ETHODUOQUAD from Akzo, DEHYQUAT SP from Henkel, and ATLAS G265 from ICI Americas.

Salts of primary, secondary, and tertiary fatty amines are also suitable cationic surfactants. The alkyl groups of such amines preferably have from about 12 to about 22 carbon atoms, and can be substituted or unsubstituted. Particularly useful are amido substituted tertiary fatty amines. Such amines, useful herein, include stearamidopropyldimethylamine, stearamidoethyldiethylamine, palmitamidopropyldimethylamine, palmitamidoethyldiethylamine, behenamidopropyldimethylamine, behenamidoethyldiethylamine, arachidamidopropyldimethylamine, arachidamidoethyldiethylamine, arachidamidoethylstearamide, diethylaminoethylstearamide. Also useful are dimethylstearamine, dimethylsoyamine, soyamine, myristylamine, tridecylamine, ethylstearylamine, N-tallowpropane diamine, ethoxylated (with 5 moles of ethylene oxide) stearylamine, dihydroxyethylstearylamine, and

arachidyl behenylamine. These amines can also be used in combination with acids such as L-glutamic acid, lactic acid, hydrochloric acid, malic acid, succinic acid, acetic acid, fumaric acid, tartaric acid, citric acid, L-glutamic hydrochloride, and mixtures thereof; more preferably L-glutamic acid, lactic acid, citric acid. Cationic amine surfactants included among those useful are disclosed in U.S. Patent 4,275,055, Nachtigal, et al., issued June 23, 1981, which is incorporated by reference herein in its entirety.

The cationic surfactants for use herein may also include a plurality of ammonium quaternary moieties or amino moieties, or a mixture thereof.

High melting point compounds

Useful hair conditioning agents include a high melting point compound having a melting point of at least about 25°C selected from the group consisting of fatty alcohols, fatty acids, fatty alcohol derivatives, fatty acid derivatives, hydrocarbons, steroids, and mixtures thereof. Without being bound by theory, it is believed that these high melting point compounds cover the hair surface and reduce friction, thereby resulting in providing smooth feel on the hair and ease of combing. It is understood by the artisan that the compounds disclosed in this section of the specification can in some instances fall into more than one classification, e.g., some fatty alcohol derivatives can also be classified as fatty acid derivatives. However, a given classification is not intended to be a limitation on that particular compound, but is done so for convenience of classification and nomenclature. Further, it is understood by the artisan that, depending on the number and position of double bonds, and length and position of the branches, certain compounds having certain required carbon atoms may have a melting point of less than about 25°C. Such compounds of low melting point are not intended to be included in this section. Nonlimiting examples of the high melting point compounds are found in International Cosmetic Ingredient Dictionary, Fifth Edition, 1993, and CTFA Cosmetic Ingredient Handbook, Second Edition, 1992, both of which are incorporated by reference herein in their entirety.

The fatty alcohols useful herein are those having from about 14 to about 30 carbon atoms, preferably from about 16 to about 22 carbon atoms. These fatty alcohols can be straight or branched chain alcohols and can be saturated or unsaturated. Nonlimiting examples of fatty alcohols include, cetyl alcohol, stearyl alcohol, behenyl alcohol, and mixtures thereof.

The fatty acids useful herein are those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and more preferably from about 16 to about 22 carbon atoms. These fatty acids can be straight or branched chain acids and can be saturated or unsaturated. Also included are diacids, triacids, and other multiple acids which meet the requirements herein. Also included herein are salts of these fatty acids. Nonlimiting examples of fatty acids include lauric acid, palmitic acid, stearic acid, behenic acid, sebacic acid, and mixtures thereof.

The fatty alcohol derivatives and fatty acid derivatives useful herein include alkyl ethers of fatty alcohols, alkoxylated fatty alcohols, alkyl ethers of alkoxylated fatty alcohols, esters of fatty alcohols, fatty acid esters of compounds having esterifiable hydroxy groups, hydroxy-substituted fatty acids, and mixtures thereof. Nonlimiting examples of fatty alcohol derivatives and fatty acid derivatives include materials such as methyl stearyl ether; the ceteth series of compounds such as ceteth-1 through ceteth-45, which are ethylene glycol ethers of cetyl alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; the steareth series of compounds such as steareth-1 through 10, which are ethylene glycol ethers of steareth alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; ceteareth 1 through ceteareth-10, which are the ethylene glycol ethers of ceteareth alcohol, i.e. a mixture of fatty alcohols containing predominantly cetyl and stearyl alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; C₁-C₃₀ alkyl ethers of the ceteth, steareth, and ceteareth compounds just described; polyoxyethylene ethers of behenyl alcohol; ethyl stearate, cetyl stearate, cetyl palmitate, stearyl stearate, myristyl myristate, polyoxyethylene cetyl ether stearate, polyoxyethylene stearyl ether stearate, polyoxyethylene lauryl ether stearate, ethyleneglycol monostearate, polyoxyethylene monostearate, polyoxyethylene distearate, propyleneglycol monostearate, propyleneglycol distearate, trimethyloipropane distearate, sorbitan stearate, polyglyceryl stearate, glyceryl monostearate, glyceryl distearate, glyceryl tristearate, and mixtures thereof.

Hydrocarbons useful herein include compounds having at least about 20 carbons.

Steroids useful herein include compounds such as cholesterol.

High melting point compounds of a single compound of high purity are preferred. Single compounds of pure fatty alcohols selected from the group of

pure cetyl alcohol, stearyl alcohol, and behenyl alcohol are highly preferred. By "pure" herein, what is meant is that the compound has a purity of at least about 90%, preferably at least about 95%. These single compounds of high purity provide good rinsability from the hair when the consumer rinses off the composition.

Commercially available high melting point compounds useful herein include: cetyl alcohol, stearyl alcohol, and behenyl alcohol having tradenames KONOL series available from New Japan Chemical (Osaka, Japan), and NAA series available from NOF (Tokyo, Japan); pure behenyl alcohol having trademark 1-DOCOSANOL available from WAKO (Osaka, Japan), various fatty acids having tradenames NEO-FAT available from Akzo (Chicago Illinois, USA), HYSTRENE available from Witco Corp. (Dublin Ohio, USA), and DERMA available from Vevy (Genova, Italy); and cholesterol having trademark NIKKOL AGUASOME LA available from Nikko. Preferably, the high melting point compound is present in the hair care compositions herein at a level of about 0.01% to about 20%.

Perfume Compounds

Perfumes in general are a desirable element of hair care compositions, since they can mask undesirable odors of other ingredients of the hair care compositions, as well as provide a pleasing aesthetic fragrance. The perfume compounds herein can additionally provide conditioning benefit.

A wide variety of perfumes are known to those skilled in the art and are commercially available. The particular perfume used is largely a matter of choice. However, the perfume should be used at a level effective for providing a noticeable aroma to the composition, or for masking undesired aroma of the composition.

In general, the compositions will comprise from about 0.01% to about 10% of a perfume compound containing carbonyl, ether or hydroxy functionalities, or mixtures thereof, preferably from about 0.05% to about 7%, more preferably from about 0.05% to about 3%. As used herein, perfume compound means aromatically active ingredients whereas "perfume" or "perfume component" includes the total perfume compounds and any accompanying perfume solvent.

Perfumes are made by those skilled in the art in a wide variety of fragrances and strengths. Typical perfumes are described in Arctander,

Perfume and Flavour Chemicals (Aroma Chemicals), Vols. I and II (1969); and Arctander, Perfume and Flavour Materials of Natural Origin (1960).

Carbonyl-containing perfume compounds include a functionality selected from the group consisting of aldehydes, ketones, esters, carboxyls, and mixtures thereof. These materials can be aliphatic (saturated or unsaturated), aromatic, heterocyclic, or alicyclic. Generally they will have from about 6 to about 18 carbon atoms.

Non-limiting examples of aliphatic aldehyde perfume compounds herein include hexyl aldehyde, heptyl aldehyde, octyl aldehyde, decyl aldehyde, undecyl aldehyde, undecylinic aldehyde, lauric aldehyde, methyl monyl acetaldehyde, myristic aldehyde, and nonyl aldehyde. Aromatic aldehydes include cinnamic aldehydes such as amyl cinnamic aldehyde and hexyl cinnamic aldehyde, anisic aldehyde, heliotropin, benzaldehyde, and vanillin. Heterocyclic aldehydes include piperonal and furan.

Non-limiting examples of aliphatic ketone perfume compounds herein include hexanone, heptanone, octanone, ionone, undecalactone, nonalactone (gomma-nonyl lactone), camphor, alpha-methyl ionone, (gomma-undecyl lactone) gomma-methyl ionone, and beta-methyl naphthyl ketone. Alicyclic ketones include, for example, amyl cyclohexanone. Aromatic ketones include beta-naphthyl methyl ketones, para-hydroxy phenolbutanone, laevo carvone, and musk ketone. Heterocyclic ketones include thujone, cedrenone, and exaltolide.

Non-limiting examples of aliphatic ester perfume compounds herein include acetates, butyrates, and formates, such as ethyl butyrate, ethyl formate, butyl acetate, isopropyl butyrate, ethyl formate, cis-3-hexenyl acetate, ethyl aceto acetate, and phenylethyl phenylacetate, carbonates such as methyl octine carbonate, and phthalates such as diethyl phthalate. Heterocyclic esters include cedryl formate and cedryl acetate. Aromatic esters include benzyl acetate, benzyl salicylate, cis-3-hexenyl acetate, cis-3-hexenyl salicylate, and methyl anthranilate. Other esters include alicyclic esters and aliphatic / alicyclic esters, such as 4 - tertiary butyl cyclohexyl acetate. Ester perfumes also include materials derived from floral materials and fruits, and from animal notes (e.g., natural isolates of civet, castoreum, and musk).

Non-limiting examples of ether perfume compounds herein include ambrox, galaxolide, and rose oxide.

Non-limiting examples of hydroxy perfume compounds herein include 1-decanol, 1-nonanol, 1-octen-3-ol, 1-terpineol (beta), 1-undecanol, 2,4-decadien-1-ol, 2,4-octadien-1-ol, 2,6-dimethyl-6-hepten-1-ol, 2-hexen-1-ol, 2-octen-1-ol, 3,7,9-trimethyl-1,6-decadien-3-ol, 3,7-dimethyl-7-methoxyoctan-2-ol, 3,7-dimethyloctan-1-ol, 3-methyloctan-3-ol, 3-octanol, 9-decen-1-ol, α -terpineol, benzyl alcohol, (+)-cedrol, cinnamic alcohol, cis-3-hexen-1-ol, citronellol, dihydrolinalool, dihydromyrcenol, dihydroterpineol, dimethyl benzyl carbinol, dimetol, ethyl linalool, eugenol, farnesol, γ -terpineol, geraniol, iso-phytol, isopropyl cyclohexanol, *t*-methanol, lavandulol, linalool, methyl phenyl carbinol, nerol, *p*-tolyl ethylalcohol, patchouli alcohol, phenyl ethyl alcohol, phenyl hexyl alcohol, roseleia, terpineol (α -form, β -form and γ -form), sandolen, tetrahydrogeraniol, tetrahydrolinalool, tetrahydromyrcenol, vanillin, ethyl-2-phenoxy ethanol, 3-methyloctan-3-ol, ethyl linalool (3,7-dimethyl-1,6-nonadien-3-ol),.

Nonlimiting examples of ether with hydroxy perfume compounds herein include 2-phenoxy ethanol, carbitol (diethylenglycol monoethylether), dipropylene glycol, and triethyl citrate.

Nonlimiting examples of ether with carbonyl perfume compounds herein include 2-phenoxyethyl-iso-butyrate, allyl cyclohexane propionate, allyl phenoxy acetate, benzyl acetate, benzyl butyrate, benzyl cinnamate, benzyl salicylate, cinnamyl cinnamate, cis-3-hexenyl acetate, cis-3-hexenyl butyrate, cis-3-hexenyl salicylate, citronellyl acetate, citronellyl formate, citronellyl propionate, dimethyl benzyl carbinyl-iso-butyrate, ethyl acetate, ethyl butyrate, ethyl caprylate, ethyl decylate, ethyl-2-methylbutyrate, ethylene brassylate, exaltolide, flor acetate, γ -decalactone, γ -dodecalactone, γ -nonalactone, γ -undecalactone, geranyl acetate, geranyl formate, helional, heliotropine, hexyl acetate, hexyl salicylate, iso-amyl acetate, iso-amyl salicylate, isopropyl myristate, linalyl acetate, menthyl acetate, methyl anthranilate, methyl benzoate, methyl phenyl carbinyl acetate, neryl acetate, phenylethyl acetate, prenyl acetate, and trichloro methyl phenyl carbinyl acetate.

In addition, certain perfume compounds may fall within one or more of the classifications described herein. As those of skill in the art will recognize, the nomenclature assigned to a particular compound will not affect its suitability for use herein.

Perfume solvents are well known in the art, and the conventional ones can be used herein, e.g., dipropylene glycol, diethylene glycol, C₁-C₆ alcohols, etc.

The hydrophobicity of the perfume raw materials, herein referred to as PRMs, may be characterized by their octanol/water partition coefficient P. The octanol/water partition coefficient of an organic compound is the ratio between its equilibrium concentration in octanol and in water. Since the partition coefficients of organic compounds are typically quite large, they are more conveniently given in the form of their logarithm to the base 10, logP.

The logP of many compounds has been reported; for example, the Pomona92 database, available from Daylight Chemical Information Systems, Inc. (Daylight CIS), contains many, along with citations to the original literature.

However, the logP values are most conveniently calculated by the "CLOGP" program, also available from Daylight CIS. This program also lists experimental logP values when they are available in the Pomona92 database. The "calculated logP" (CLogP) is determined by the fragment approach of Hansch and Leo (A. Leo, Comprehensive Medicinal Chemistry, Vol. 4; C. Hansch, P.G. Sammens, J.B. Taylor and C.A. Ramden, Eds., p. 295, Pergamon Press, 1990). The fragment approach is based on the chemical structure of a compound and takes into account the number and type of atoms, the atom connectivity, and chemical bonding. The CLogP values, which are the most reliable and widely used estimates for this physiochemical property, can be used instead of the experimental logP values in the selection of perfume compounds. The perfume compounds of the present compositions preferably have a CLogP value of between -1.0 and 8.0.

Without being bound by theory, it is believed that the addition of perfume compounds to the compositions of the present invention may affect the stability, including viscosity and phase separation, and the conditioning properties of the composition.

For shampoo compositions, viscosity increase of the composition may be observed by the addition of PRMs. It is believed that the viscosity increase is caused by solubilization of the PRMs into the palisade layer between hydrocarbon core and micelle surface. These molecules expand the area between surfactant molecules in the micelle and the micelle size increases. Enlarged micelle increases an intermicelle interaction, such as entanglement of

the cylindrical micelle, as a result of increasing occupied volume fraction of the micelle.

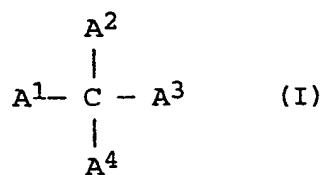
Water-insoluble High Molecular Weight Oily Compound

The water-insoluble high molecular weight oily compounds of the present invention are those which provide excellent conditioning benefits such as smoothness to the hair and ease of combing. Without being bound by theory, it is believed that, the water-insoluble high molecular weight oily compound of this invention is capable of being deposited on and conditions the hair. It is also believed that this water-insoluble high molecular weight oily compound covers the surface of the hair and as a result reduces hair friction to deliver smoothness to the hair. The water-insoluble high molecular weight oily compound is chemically stable under normal use and storage conditions.

As used herein, the term "water-insoluble oily compound" is meant that the compound is not sufficiently soluble in water at room temperature. When the compound is mixed with water at above 1.0% concentration, more preferably at above 0.5% concentration, the compound is temporarily dispersed to form unstable colloid in water, then is quickly separated from water into two phases in a short period.

The water-insoluble high molecular weight oily compound useful in the present invention has a molecular weight of at least about 800, preferably at least about 1000, more preferably at least about 1200 as long as the compound has a specific gravity of at least about 0.9, and is in a liquid form at 25°C.

The water-insoluble high molecular weight oily compound useful herein include those of the following formula (I):



wherein A¹, A², A³ and A⁴ are independently alkyl, alkenyl, aryl, alkylaryl, hydroxyalkyl, alkoxy, alkoxyalkyl, acyl, acylalkyl, and alkylacyloxy group having C₁ to about C₃₀ or the formula -(CH₂)_n-O-OCR wherein R is from C₁ to about C₃₀ of branched or straight chain alkyl or alkenyl and n is an integer from 1 to about 30, and preferably equals 1.

Preferably, the water-insoluble high molecular weight oily compound of this invention is an ester wherein all of A¹, A², A³ and A⁴ are ester groups or

three of A¹, A², A³ and A⁴ are ester groups and the remainder is an alkyl group. More preferably, the water-insoluble high molecular weight oily compounds of this invention are esters of a fatty acid of from about C₁₂ to about C₂₂ with pentaerythritol, esters of a fatty acid of from about C₁₂ to about C₂₂ with trimethylolalkane, and mixtures thereof.

Preferable water-insoluble high molecular weight oily compounds are those which have the same substituent for A¹, A², A³ and A⁴ and wherein these substituents are ester groups.

The preferred water-insoluble high molecular weight oily compounds of this invention are pentaerythritol tetraisostearate, trimethylolpropane triisostearate, pentaerythritol tetraoleate, trimethylolpropane trioleate, and mixtures thereof.

Suitable ester compounds include, for example, pentaerythritol tetraisostearate and trimethylolpropane triisostearate which are available from Kokyo Alcohol with tradenames KAKPTI and KAKTTI, and pentaerythritol tetraoleate and trimethylolpropane trioleate which are available from Shinnihon Rika with tradenames PTO and ENUJERUBU series, respectively.

The water-insoluble high molecular weight oily compound is used at a level of from about 0.1% to about 20.0%, preferably from about 0.1% to about 10.0%, more preferably from about 0.2% to about 5.0% by weight of the composition.

DETERSIVE SURFACTANT

Compositions herein may include a detergents surfactant. The detergents surfactants herein are those suitable for cleansing the hair. Detergents surfactants useful herein include anionic surfactants, amphoteric and zwitterionic surfactants, and nonionic surfactants.

The detergents surfactants, when present, are preferably included at a level of from about 0.01% to about 75% by weight of the composition. Two or more surfactants can be used.

Anionic Surfactant

Anionic surfactants useful herein include alkyl and alkyl ether sulfates. These materials have the respective formulae ROSO_3M and $\text{RO}(\text{C}_2\text{H}_4\text{O})_x\text{SO}_3\text{M}$, wherein R is alkyl or alkenyl of from about 8 to about 30 carbon atoms, x is 1 to about 10, and M is hydrogen or a cation such as ammonium, alkanolammonium (e.g., triethanolammonium), a monovalent metal cation (e.g., sodium and potassium), or a polyvalent metal cation (e.g., magnesium and calcium). Preferably, M should be chosen such that the anionic surfactant component is water soluble. The anionic surfactant or surfactants should be chosen such that the Krafft temperature is about 15°C or less, preferably about 10°C or less, and more preferably about 0°C or less. It is also preferred that the anionic surfactant be soluble in the composition hereof.

Krafft temperature refers to the point at which solubility of an ionic surfactant becomes determined by crystal lattice energy and heat of hydration, and corresponds to a point at which solubility undergoes a sharp, discontinuous increase with increasing temperature. Each type of surfactant will have its own characteristic Krafft temperature. Krafft temperature for ionic surfactants is, in general, well known and understood in the art. See, for example, Myers, Drew, Surfactant Science and Technology, pp. 82-85, VCH Publishers, Inc. (New York, New York, USA), 1988 (ISBN 0-89573-399-0), which is incorporated by reference herein in its entirety.

In the alkyl and alkyl ether sulfates described above, preferably R has from about 8 to about 18 carbon atoms in both the alkyl and alkyl ether sulfates. The alkyl ether sulfates are typically made as condensation products of ethylene oxide and monohydric alcohols having from about 8 to about 24 carbon atoms. The alcohols can be derived from fats, e.g., coconut oil, palm oil, tallow, or the like, or the alcohols can be synthetic. Lauryl alcohol and straight chain alcohols derived from coconut oil and palm oil are preferred herein. Such alcohols are reacted with 1 to about 10, and especially about 3, molar proportions of ethylene oxide and the resulting mixture of molecular species having, for example, an average of 3 moles of ethylene oxide per mole of alcohol, is sulfated and neutralized.

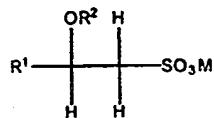
Specific examples of alkyl ether sulfates which can be used are sodium and ammonium salts of coconut alkyl triethylene glycol ether sulfate; tallow alkyl triethylene glycol ether sulfate, and tallow alkyl hexaoxyethylene sulfate. Highly preferred alkyl ether sulfates are those comprising a mixture of individual

compounds, said mixture having an average alkyl chain length of from about 8 to about 16 carbon atoms and an average degree of ethoxylation of from 1 to about 4 moles of ethylene oxide. Such a mixture also comprises from 0% to about 20% by weight C₁₂₋₁₃ compounds; from about 60% to about 100% by weight of C₁₄₋₁₅₋₁₆ compounds, from 0% to about 20% by weight of C₁₇₋₁₈₋₁₉ compounds; from about 3% to about 30% by weight of compounds having a degree of ethoxylation of 0; from about 45% to about 90% by weight of compounds having a degree of ethoxylation of from 1 to about 4; from about 10% to about 25% by weight of compounds having a degree of ethoxylation of from about 4 to about 8; and from about 0.1% to about 15% by weight of compounds having a degree of ethoxylation greater than about 8.

Other suitable anionic surfactants are the water-soluble salts of organic, sulfuric acid reaction products of the general formula [R¹-SO₃-M] where R¹ is selected from the group consisting of a straight or branched chain, saturated aliphatic hydrocarbon radical having from about 8 to about 24, preferably about 8 to about 18, carbon atoms; and M is as previously described above in this section. Examples of such surfactants are the salts of an organic sulfuric acid reaction product of a hydrocarbon of the methane series, including iso-, neo-, and n-paraffins, having about 8 to about 24 carbon atoms, preferably about 8 to about 18 carbon atoms and a sulfonating agent, e.g., SO₃, H₂SO₄, obtained according to known sulfonation methods, including bleaching and hydrolysis. Preferred are alkali metal and ammonium sulfonated C₈₋₁₈ n-paraffins.

Still other suitable anionic surfactants are the reaction products of fatty acids esterified with isethionic acid and neutralized with sodium hydroxide where, for example, the fatty acids are derived from coconut or palm oil; or sodium or potassium salts of fatty acid amides of methyl tauride in which the fatty acids, for example, are derived from coconut oil. Other similar anionic surfactants are described in U.S. Patents 2,486,921, 2,486,922, and 2,396,278, which are incorporated by reference herein in their entirety.

Another class of anionic surfactants suitable for use in the shampoo compositions are the β -alkyloxy alkane sulfonates. These compounds have the following formula:

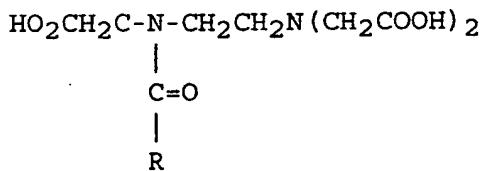


where R¹ is a straight chain alkyl group having from about 6 to about 20 carbon atoms, R² is a lower alkyl group having from about 1, preferred, to about 3 carbon atoms, and M is as hereinbefore described. Many other anionic surfactants suitable for use in the shampoo compositions are described in McCutcheon's, Emulsifiers and Detergents, 1989 Annual, published by M. C. Publishing Co., and in U.S. Patent 3,929,678, which descriptions are incorporated herein by reference in their entirety. Preferred anionic surfactants for use in the shampoo compositions include ammonium lauryl sulfate, ammonium laureth sulfate, triethylamine lauryl sulfate, triethylamine laureth sulfate, triethanolamine lauryl sulfate, triethanolamine laureth sulfate, monoethanolamine lauryl sulfate, monoethanolamine laureth sulfate, diethanolamine lauryl sulfate, diethanolamine laureth sulfate, lauric monoglyceride sodium sulfate, sodium lauryl sulfate, sodium laureth sulfate, potassium lauryl sulfate, potassium laureth sulfate, ammonium cocoyl sulfate, ammonium lauroyl sulfate, sodium cocoyl sulfate, sodium lauroyl sulfate, potassium cocoyl sulfate, potassium lauryl sulfate, triethanolamine lauryl sulfate, triethanolamine lauryl sulfate, monoethanolamine cocoyl sulfate, monoethanolamine lauryl sulfate, sodium tridecyl benzene sulfonate, and sodium dodecyl benzene sulfonate, and mixtures thereof.

Other anionic surfactants for use herein include polyhydrophilic anionic surfactants. By "polyhydrophilic" herein, is meant a surfactant that has at least two hydrophilic groups which provide a hydrophilic nature. Polyhydrophilic surfactants useful herein are only those having at least two hydrophilic groups in the molecule, and is not intended to encompass those which only have one hydrophilic group. One molecule of the polyhydrophilic anionic surfactant herein may comprise the same hydrophilic groups, or different hydrophilic groups. Specifically, the polyhydrophilic anionic surfactants comprise at least one group selected from the group consisting of carboxy, hydroxy, sulfate, sulfonate, and phosphate. Suitable polyhydrophilic anionic surfactants are those which comprise at least one of a carboxy, sulfate, or sulfonate group, more preferably those which comprise at least one carboxy group.

Nonlimiting examples of polyhydrophilic anionic surfactants include N-acyl-L-glutamates such as N-cocoyl-L-glutamate and, N-lauroyl-L-glutamate, laurimino dipropionate, N-acyl-L-aspartate, di-(N-lauroyl N-methyl taurate), polyoxyethylene laurylsulfosuccinate, disodium N-octadecylsulfosuccinate; disodium lauryl sulfosuccinate; diammonium lauryl sulfosuccinate; tetra sodium

N-(1,2-dicarboxyethyl)-N-octadecylsulfosuccinate; the diamyl ester of sodium sulfosuccinic acid; the dihexyl ester of sodium sulfosuccinic acid; and the dioctyl ester of sodium sulfosuccinic acid, and 2-cocoalkyl N-carboxyethyl N-carboxyethoxyethyl imidazolinium betaine, lauroamphohydroxypropylsulfonate, cocoglyceryl ether salts, cocoglyceride sulfate, lauroyl isethionate, lauroamphoacetate, and those of the following formula:



wherein R is an alkyl of 8 to 18 carbons. Other polyhydrophilic anionic surfactants include olefin sulfonates having about 10 to about 24 carbon atoms. The term "olefin sulfonates" is used herein to mean compounds which can be produced by the sulfonation of alpha-olefins by means of uncomplexed sulfur trioxide, followed by neutralization of the acid reaction mixture in conditions such that any sulfones which have been formed in the reaction are hydrolyzed to give the corresponding hydroxy-alkanesulfonates. The sulfur trioxide can be liquid or gaseous, and is usually, but not necessarily, diluted by inert diluents, for example by liquid SO_2 , chlorinated hydrocarbons, etc., when used in the liquid form, or by air, nitrogen, gaseous SO_2 , etc., when used in the gaseous form. The α -olefins from which the olefin sulfonates are derived are mono-olefins having about 8 to about 24 carbon atoms, preferably about 10 to about 16 carbon atoms. Preferably, they are straight chain olefins. In addition to the true alkene sulfonates and a proportion of hydroxy-alkanesulfonates, the olefin sulfonates can contain minor amounts of other materials, such as alkene disulfonates depending upon the reaction conditions, proportion of reactants, the nature of the starting olefins and impurities in the olefin stock and side reactions during the sulfonation process. A specific α -olefin sulfonate mixture of the above type is described more fully in U.S. Patent 3,332,880, to Pflaumer and Kessler, issued July 25, 1967, which is incorporated by reference herein in its entirety.

Another class of polyhydrophilic anionic surfactants are amino acid surfactants which are surfactants that have the basic chemical structure of an amino acid compound, i.e., that contains a structural component of one of the naturally-occurring amino acids. It is understood by the artisan that some

surfactants may be regarded as both a polyhydrophilic anionic surfactant, and an amino acid surfactant. These surfactants are suitable anionic surfactants.

Nonlimiting examples of amino acid surfactants include, N-cocoylalaninate, N-acyl-N-methyl- β -alanate, N-acylsarcosinate; N-alkylamino propionates and N-alkyliminodipropionates, specific examples of which include N-lauryl- β -amino propionic acid or salts thereof, and N-lauryl- β -imino-dipropionate, N-acyl-DL-alaninate, sodium lauryl sarcosinate, sodium lauroyl sarcosinate, lauryl sarcosine, cocoyl sarcosine, N-acyl-N-methyl taurate, lauroyl taurate, and lauroyl lactylate.

Commercially available anionic surfactants suitable are N-acyl-L-glutamate with a tradename AMISOFT CT-12S, N-acyl potassiumglycine with a tradename AMILITE GCK-12, lauroyl glutamate with a tradename AMISOFT LS-11, and N-acyl-DL-alaninate with tradename AMILITE ACT12 supplied by Ajinomoto; acylaspartate with tradenames ASPARACK and AAS supplied by Mitsubishi Chemical; and acyl derivaties of tradename ED3A supplied by Hampshire Chemical Corp.

Optionally the counter ion of anionic surfactants may be polyvalent cations. It has been found that these anionic surfactants, along with the cationic conditioning agents, and polyvalent metal cations as described later, form a coacervate in the compositions. Cationic conditioning agents may be included in the present composition to provide a shampoo which both cleanse and condition the hair from a single product.

Coacervate formulation is dependent upon a variety of criteria such as molecular weight, component concentration, and ratio of interacting ionic components ionic strength, charge density of the cationic and anionic components, pH, and temperature. Coacervate systems and the effect of these parameters are known in the art.

It is believed to be particularly advantageous, for the anionic surfactants and the polyvalent metal cations at certain levels to be present with the cationic conditioning agents in a coacervate phase. The coacervates formed in the compositions are believed to readily deposit on the hair upon diluting the coacervate with abundant water, i.e., rinsing of the shampoo.

Without being bound by theory, it is believed that the coacervates provide two major effects to the present shampoo composition. First, it reduces the Critical Micelle Concentration (hereinafter "CMC") of the composition. The reduction of the CMC relates to reduction of the surface tension, thereby

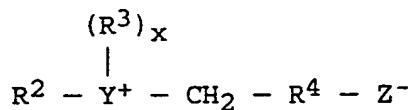
improving lather performance. Second, the existence of the anionic surfactants along with the polyvalent metal cations expand the coacervate region in the composition. As the cationic conditioning agents in the composition are mainly delivered to the hair via these coacervates, expansion of the coacervate region results in delivery of more cationic conditioning agents to the hair. Consequently, compositions which both cleanse and condition the hair from a single product, which have improved overall conditioning benefits and improved lathering are provided.

Techniques of analysis of formation of complex coacervates are known in the art. For example, microscopic analysis of the shampoo compositions, at any chosen stage of dilution, can be utilized to identify whether a coacervate phase has formed. Such coacervate phase will be identifiable as an additional emulsified phase in the composition. The use of dyes can aid in distinguishing the coacervate phase from other insoluble phases dispersed in the shampoo composition.

Amphoteric And Zwitterionic Surfactant

Amphoteric surfactants for use herein include the derivatives of aliphatic secondary and tertiary amines in which the aliphatic radical is straight or branched and one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic water solubilizing group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate.

Zwitterionic surfactants for use herein include the derivatives of aliphatic quaternary ammonium, phosphonium, and sulfonium compounds, in which the aliphatic radicals are straight or branched, and wherein one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate. A general formula for these compounds is:

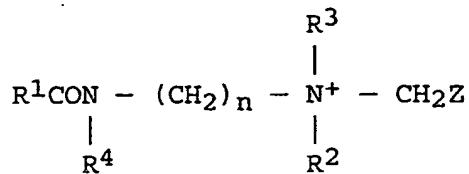


where R^2 contains an alkyl, alkenyl, or hydroxy alkyl radical of from about 8 to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties and from 0 to about 1 glyceryl moiety; Y is selected from the group consisting of nitrogen, phosphorus, and sulfur atoms; R^3 is an alkyl or monohydroxyalkyl group containing 1 to about 3 carbon atoms; X is 1 when Y is a sulfur atom, and 2 when Y is a nitrogen or phosphorus atom; R^4 is an alkylene or hydroxyalkylene of from 1 to about 4 carbon atoms and Z is a radical selected from the group consisting of carboxylate, sulfonate, sulfate, phosphonate, and phosphate groups.

Examples of amphoteric and zwitterionic surfactants also include sultaines and amidosultaines. Sultaines, including amidosultaines, include for example, cocodimethylpropylsultaine, stearyldimethylpropylsultaine, lauryl-bis-(2-hydroxyethyl)propylsultaine and the like; and the amidosultaines such as cocamidodimethylpropylsultaine, stearylamidododimethylpropylsultaine, laurylamido-bis-(2-hydroxyethyl)propylsultaine, and the like. Preferred are amidohydroxysultaines such as the C₈-C₁₈ hydrocarbylamidopropyl hydroxysultaines, especially C₈-C₁₄ hydrocarbylamidopropylhydroxysultaines, e.g., laurylamidopropylhydroxysultaine and cocamidopropylhydroxysultaine. Other sultaines are described in U.S. Patent 3,950,417, which is incorporated herein by reference in its entirety.

Other suitable amphoteric surfactants are the aminoalkanoates of the formula $RNH(CH_2)_nCOOM$, the iminodialkanoates of the formula $RN[(CH_2)_mCOOM]_2$ and mixtures thereof; wherein n and m are numbers from 1 to about 4, R is C₈ - C₂₂ alkyl or alkenyl, and M is hydrogen, alkali metal, alkaline earth metal, ammonium or alkanolammonium.

Other suitable amphoteric surfactants include those represented by the formula :

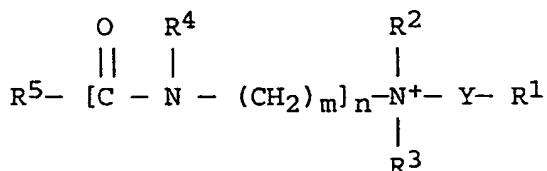


wherein R¹ is C₈ - C₂₂ alkyl or alkenyl, preferably C₈ - C₁₆, R² and R³ is independently selected from the group consisting of hydrogen, -CH₂CO₂M, -CH₂CH₂OH, -CH₂CH₂OCH₂CH₂COOM, or -(CH₂CH₂O)_mH wherein m is an integer from 1 to about 25, and R⁴ is hydrogen, -CH₂CH₂OH, or CH₂CH₂OCH₂CH₂COOM, Z is CO₂M or CH₂CO₂M, n is 2 or 3, preferably 2, M is hydrogen or a cation, such as alkali metal (e.g., lithium, sodium, potassium), alkaline earth metal (beryllium, magnesium, calcium, strontium, barium), or ammonium. This type of surfactant is sometimes classified as an imidazoline-type amphoteric surfactant, although it should be recognized that it does not necessarily have to be derived, directly or indirectly, through an imidazoline intermediate. Suitable materials of this type are marketed under the tradename MIRANOL and are understood to comprise a complex mixture of species, and can exist in protonated and non-protonated species depending upon pH with respect to species that can have a hydrogen at R². All such variations and species are meant to be encompassed by the above formula.

Examples of surfactants of the above formula are monocarboxylates and di-carboxylates. Examples of these materials include cocoamphocarboxypropionate, cocoamphocarboxypropionic acid, cocoamphocarboxyglycinate (alternately referred to as cocoamphodiacetate), and cocoamphoacetate.

Commercial amphoteric surfactants include those sold under the trade names MIRANOL C2M CONC. N.P., MIRANOL C2M CONC. O.P., MIRANOL C2M SF, MIRANOL CM SPECIAL (Miranol, Inc.); ALKATERIC 2CIB (Alkaril Chemicals); AMPHOTERGE W-2 (Lonza, Inc.); MONATERIC CDX-38, MONATERIC CSH-32 (Mona Industries); REWOTERIC AM-2C (Rewo Chemical Group); and SCHERCOTERIC MS-2 (Scher Chemicals).

Betaine surfactants, i.e. zwitterionic surfactants, suitable for use in the conditioning compositions are those represented by the formula:



wherein: R¹ is a member selected from the group consisting of COOM and CH(OH)CH₂SO₃M

R^2 is lower alkyl or hydroxyalkyl; R^3 is lower alkyl or hydroxyalkyl; R^4 is a member selected from the group consisting of hydrogen and lower alkyl; R^5 is higher alkyl or alkenyl; Y is lower alkyl, preferably methyl; m is an integer from 2 to 7, preferably from 2 to 3; n is the integer 1 or 0; M is hydrogen or a cation, as previously described, such as an alkali metal, alkaline earth metal, or ammonium. The term "lower alkyl" or "hydroxyalkyl" means straight or branch chained, saturated, aliphatic hydrocarbon radicals and substituted hydrocarbon radicals having from one to about three carbon atoms such as, for example, methyl, ethyl, propyl, isopropyl, hydroxypropyl, hydroxyethyl, and the like. The term "higher alkyl or alkenyl" means straight or branch chained saturated (i.e., "higher alkyl") and unsaturated (i.e., "higher alkenyl") aliphatic hydrocarbon radicals having from about 8 to about 20 carbon atoms such as, for example, lauryl, cetyl, stearyl, oleyl, and the like. It should be understood that the term "higher alkyl or alkenyl" includes mixtures of radicals which may contain one or more intermediate linkages such as ether or polyether linkages or non-functional substituents such as hydroxyl or halogen radicals wherein the radical remains of hydrophobic character.

Examples of surfactant betaines of the above formula wherein n is zero which are useful herein include the alkylbetaines such as cocodimethylcarboxymethylbetaine, lauryldimethylcarboxymethylbetaine, lauryldimethyl- α -carboxyethylbetaine, cetyltrimethylcarboxymethylbetaine, lauryl-bis-(2-hydroxyethyl)-carboxymethylbetaine, stearyl-bis-(2-hydroxypropyl)carboxymethylbetaine, oleyldimethyl- γ -carboxypropylbetaine, lauryl-bis-(2-hydroxypropyl)- α -carboxyethylbetaine, etc. The sulfobetaines may be represented by cocodimethylsulfopropylbetaine, stearyltrimethylsulfopropyl betaine, lauryl-bis-(2-hydroxyethyl)-sulfopropylbetaine, and the like.

Specific examples of amido betaines and amidosulfobetaines useful in the conditioning compositions include the amidocarboxybetaines, such as cocamido dimethylcarboxymethylbetaine, laurylamidodimethylcarboxymethylbetaine, cetylamidodimethylcarboxymethylbetaine, laurylamido-bis-(2-hydroxyethyl)carboxymethylbetaine, cocamido-bis-(2-hydroxyethyl)-carboxymethylbetaine, etc. The amidosulfobetaines may be represented by cocamidodimethyl sulfopropylbetaine, stearyltrimodimethylsulfopropylbetaine, laurylamido-bis-(2-hydroxyethyl)-sulfopropylbetaine, and the like.

Nonionic Surfactant

The compositions of the present invention can comprise a nonionic surfactant. Nonionic surfactants include those compounds produced by condensation of alkylene oxide groups, hydrophilic in nature, with an organic hydrophobic compound, which may be aliphatic or alkyl aromatic in nature.

Preferred nonlimiting examples of nonionic surfactants for use in the shampoo compositions include the following:

(1) polyethylene oxide condensates of alkyl phenols, e.g., the condensation products of alkyl phenols having an alkyl group containing from about 6 to about 20 carbon atoms in either a straight chain or branched chain configuration, with ethylene oxide, the said ethylene oxide being present in amounts equal to from about 10 to about 60 moles of ethylene oxide per mole of alkyl phenol;

(2) those derived from the condensation of ethylene oxide with the product resulting from the reaction of propylene oxide and ethylene diamine products;

(3) condensation products of aliphatic alcohols having from about 8 to about 18 carbon atoms, in either straight chain or branched chain configurations, with ethylene oxide, e.g., a coconut alcohol ethylene oxide condensate having from about 10 to about 30 moles of ethylene oxide per mole of coconut alcohol, the coconut alcohol fraction having from about 8 to about 14 carbon atoms;

(4) long chain tertiary amine oxides of the formula [$R^1R^2R^3N \rightarrow O$] where R^1 contains an alkyl, alkenyl or monohydroxy alkyl radical of from about 8 to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties, and from 0 to about 1 glycerol moiety, and R^2 and R^3 contain from about 1 to about 3 carbon atoms and from 0 to about 1 hydroxy group, e.g., methyl, ethyl, propyl, hydroxyethyl, or hydroxypropyl radicals;

(5) long chain tertiary phosphine oxides of the formula [$RR'R''P \rightarrow O$] where R contains an alkyl, alkenyl or monohydroxyalkyl radical ranging from about 8 to about 18 carbon atoms in chain length, from 0 to about 10 ethylene oxide moieties and from 0 to 1 glycerol moieties and R' and R'' are each alkyl or monohydroxyalkyl groups containing from about 1 to about 3 carbon atoms;

(6) long chain dialkyl sulfoxides containing one short chain alkyl or hydroxy alkyl radical of from 1 to about 3 carbon atoms (usually methyl) and one long hydrophobic chain which include alkyl, alkenyl, hydroxy alkyl, or keto alkyl

radicals containing from about 8 to about 20 carbon atoms, from 0 to about 10 ethylene oxide moieties and from 0 to 1 glyceryl moieties;

(7) alkyl polysaccharide (APS) surfactants (e.g. alkyl polyglycosides), examples of which are described in U.S. Patent 4,565,647, which is incorporated herein by reference in its entirety, and which discloses APS surfactants having a hydrophobic group with about 6 to about 30 carbon atoms and a polysaccharide (e.g., polyglycoside) as the hydrophilic group; optionally, there can be a polyalkylene-oxide group joining the hydrophobic and hydrophilic moieties; and the alkyl group (i.e., the hydrophobic moiety) can be saturated or unsaturated, branched or unbranched, and unsubstituted or substituted (e.g., with hydroxy or cyclic rings); a preferred material is alkyl polyglucoside which is commercially available from Henkel, ICI Americas, and Seppic; and

(8) polyoxyethylene alkyl ethers such as those of the formula $RO(CH_2CH_2)_nH$ and polyethylene glycol (PEG) glycetyl fatty esters, such as those of the formula $R(O)OCH_2CH(OH)CH_2(OCH_2CH_2)_nOH$, wherein n is from 1 to about 200, preferably from about 20 to about 100, and R is an alkyl having from about 8 to about 22 carbon atoms.

CARRIER

The hair care composition of the present invention may comprise a carrier. The level and species of the carrier are selected according to the compatibility with other components, and desired characteristic of the product. For example, a high percentage of volatile solvents of low boiling point and/or propellant are suitably used for product forms aimed to be left on the hair. On the other hand, water solutions of volatile and non-volatile solvents are suitably used for product forms aimed to be rinsed off the hair after washing or treating the hair with the product.

The carrier useful in the present invention include volatile solvents, non-volatile solvents, propellants, and mixtures thereof.

Volatile solvents useful herein include water, lower alkyl alcohols having from 1 to 3 carbons, and hydrocarbons having from about 5 to about 8 carbons. The preferred volatile solvents are water, ethanol, isopropanol, pentane, hexane, and heptane. The water useful herein include deionized water and water from natural sources containing mineral cations. Deionized water is preferred.

Non-volatile solvents useful herein include alkyl alcohols having more than 3 carbons, and polyhydric alcohols. The polyhydric alcohols useful herein

include 1,2-propane diol or propylene glycol, 1,3-propane diol, hexylene glycol, glycerin, diethylene glycol, dipropylene glycol, 1,2-butyleneglycol, and 1,4-butyleneglycol.

Propellants may be used for mousse and hair spray product forms. Propellants, when used in the present invention, are selected depending on variables such as the remainder of components, the package, and whether the product is designed to be used standing or invert.

Propellants useful herein include fluorohydrocarbons such as difluoroethane 152a (supplied by DuPont), dimethylether, and hydrocarbons such as propane, isobutane, n-butane, mixtures of hydrocarbons such as LPG (liquefied petroleum gas), carbon dioxide, nitrous oxide, nitrogen, and compressed air.

ADDITIONAL CONDITIONING AGENTS

Additional conditioning agents useful herein include oily compounds and nonionic polymers.

Oily Compound

Oily compound useful herein have a melting point of not more than about 25°C selected from the group consisting of a first oily compound, a second oily compound, and mixtures thereof. The oily compounds useful herein may be volatile or nonvolatile. Without being bound by theory, it is believed that, the oily compounds may penetrate the hair to modify the hydroxy bonds of the hair, thereby resulting in providing softness and flexibility to the hair. The oily compound may comprise either the first oily compound or the second oily compound as described herein. Preferably, a mixture of the first oily compound and the second oily compound is used. The oily compounds of this section are to be distinguished from the high melting point compounds described above. Nonlimiting examples of the oily compounds are found in International Cosmetic Ingredient Dictionary, Fifth Edition, 1993, and CTFA Cosmetic Ingredient Handbook, Second Edition, 1992, both of which are incorporated by reference herein in their entirety.

First Oily Compound

The fatty alcohols useful herein include those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and more preferably from about 16 to about 22 carbon atoms. These fatty alcohols can be straight or branched chain alcohols and can be saturated or unsaturated alcohols, preferably unsaturated alcohols. Nonlimiting examples of these

compounds include oleyl alcohol, palmitoleic alcohol, isostearyl alcohol, isocetyl alcohol, undecanol, octyl dodecanol, octyl decanol, octyl alcohol, caprylic alcohol, decyl alcohol and lauryl alcohol.

The fatty acids useful herein include those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and more preferably from about 16 to about 22 carbon atoms. These fatty acids can be straight or branched chain acids and can be saturated or unsaturated. Suitable fatty acids include, for example, oleic acid, linoleic acid, isostearic acid, linolenic acid, ethyl linolenic acid, ethyl linolenic acid, arachidonic acid, and ricinolic acid.

The fatty acid derivatives and fatty alcohol derivatives are defined herein to include, for example, esters of fatty alcohols, alkoxylated fatty alcohols, alkyl ethers of fatty alcohols, alkyl ethers of alkoxylated fatty alcohols, and mixtures thereof. Nonlimiting examples of fatty acid derivatives and fatty alcohol derivatives, include, for example, methyl linoleate, ethyl linoleate, isopropyl linoleate, isodecyl oleate, isopropyl oleate, ethyl oleate, octyldodecyl oleate, oleyl oleate, decyl oleate, butyl oleate, methyl oleate, octyldodecyl stearate, octyldodecyl isostearate, octyldodecyl isopalmitate, octyl isopelargonate, octyl pelargonate, hexyl isostearate, isopropyl isostearate, isodecyl isononanoate, Oleth-2, pentaerythritol tetraoleate, pentaerythritol tetraisostearate, trimethylolpropane trioleate, and trimethylolpropane triisostearate.

Commercially available first oily compounds useful herein include: oleyl alcohol with tradename UNJECOL 90BHR available from New Japan Chemical, pentaerythritol tetraisostearate and trimethylolpropane triisostearate with tradenames KAKPTI and KAKTTI available from Kokyu Alcohol (Chiba, Japan), pentaerythritol tetraoleate having the same tradename as the compound name available from New Japan Chemical, trimethylolpropane trioleate with a tradename ENUJERUBU series available from New Japan Chemical, various liquid esters with tradenames SCHERCEMOL series available from Scher, and hexyl isostearate with a tradename HIS and isopropyl isostearate having a tradename ZPIS available from Kokyu Alcohol.

Second Oily Compound

The second oily compounds useful herein include straight chain, cyclic, and branched chain hydrocarbons which can be either saturated or unsaturated, so long as they have a melting point of not more than about 25°C. These hydrocarbons have from about 12 to about 40 carbon atoms, preferably from about 12 to about 30 carbon atoms, and preferably from about 12 to about 22

carbon atoms. Also encompassed herein are polymeric hydrocarbons of alkenyl monomers, such as polymers of C2-6 alkenyl monomers. These polymers can be straight or branched chain polymers. The straight chain polymers will typically be relatively short in length, having a total number of carbon atoms as described above. The branched chain polymers can have substantially higher chain lengths. The number average molecular weight of such materials can vary widely, but will typically be up to about 500, preferably from about 200 to about 400, and more preferably from about 300 to about 350. Also useful herein are the various grades of mineral oils. Mineral oils are liquid mixtures of hydrocarbons that are obtained from petroleum. Specific examples of suitable hydrocarbon materials include paraffin oil, mineral oil, dodecane, isododecane, hexadecane, isohexadecane, eicosene, isoeicosene, tridecane, tetradecane, polybutene, polyisobutene, and mixtures thereof. Preferred for use herein are hydrocarbons selected from the group consisting of mineral oil, isododecane, isohexadecane, polybutene, polyisobutene, and mixtures thereof.

Commercially available second oily compounds useful herein include isododecane, isohexadecane, and isoeicosene with tradenames PERMETHYL 99A, PERMETHYL 101A, and PERMETHYL 1082, available from Presperse (South Plainfield New Jersey, USA), a copolymer of isobutene and normal butene with tradenames INDOPOL H-100 available from Amoco Chemicals (Chicago Illinois, USA), mineral oil with tradename BENOL available from Witco, isoparaffin with tradename ISOPAR from Exxon Chemical Co. (Houston Texas, USA), α -olefin oligomer with tradename PURESYN 6 from Mobil Chemical Co., and trimethylolpropane tricaprylate/tricaprate with tradename MOBIL ESTER P43 from Mobil Chemical Co.

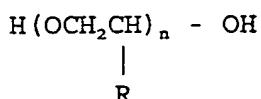
Nonionic Polymer

Nonionic polymers useful herein include cellulose derivatives, hydrophobically modified cellulose derivatives, ethylene oxide polymers, and ethylene oxide/propylene oxide based polymers. Suitable nonionic polymers are cellulose derivatives including methylcellulose with tradename BENECEL, hydroxyethyl cellulose with tradename NATROSOL, hydroxypropyl cellulose with tradename KLUCEL, cetyl hydroxyethyl cellulose with tradename POLYSURF 67, all supplied by Hercules. Other suitable nonionic polymers are ethylene oxide and/or propylene oxide based polymers with tradenames CARBOWAX PEGs, POLYOX WASRs, and UCON FLUIDS, all supplied by Amerchol.

Polyalkylene Glycols

These compounds are particularly useful for compositions which are designed to impart a soft, moist feeling to the hair. When present, the polyalkylene glycol is typically used at a level from about 0.025% to about 1.5%, preferably from about 0.05% to about 1%, and more preferably from about 0.1% to about 0.5% of the compositions.

The polyalkylene glycols are characterized by the general formula:



wherein R is selected from the group consisting of H, methyl, and mixtures thereof. When R is H, these materials are polymers of ethylene oxide, which are also known as polyethylene oxides, polyoxyethylenes, and polyethylene glycols. When R is methyl, these materials are polymers of propylene oxide, which are also known as polypropylene oxides, polyoxypropylenes, and polypropylene glycols. When R is methyl, it is also understood that various positional isomers of the resulting polymers can exist.

In the above structure, n has an average value of from about 1500 to about 25,000, preferably from about 2500 to about 20,000, and more preferably from about 3500 to about 15,000.

Polyethylene glycol polymers useful herein are PEG-2M wherein R equals H and n has an average value of about 2,000 (PEG-2M is also known as Polyox WSR® N-10, which is available from Union Carbide and as PEG-2,000); PEG-5M wherein R equals H and n has an average value of about 5,000 (PEG-5M is also known as Polyox WSR® N-35 and Polyox WSR® N-80, both available from Union Carbide and as PEG-5,000 and Polyethylene Glycol 300,000); PEG-7M wherein R equals H and n has an average value of about 7,000 (PEG-7M is also known as Polyox WSR® N-750 available from Union Carbide); PEG-9M wherein R equals H and n has an average value of about 9,000 (PEG 9-M is also known as Polyox WSR® N-3333 available from Union Carbide); and PEG-14 M wherein R equals H and n has an average value of about 14,000 (PEG-14M is also known as Polyox WSR® N-3000 available from Union Carbide).

Other useful polymers include the polypropylene glycols and mixed polyethylene/polypropylene glycols.

ADDITIONAL COMPONENTS

The compositions of the present invention may include a variety of additional components, which may be selected by the artisan according to the desired characteristics of the final product. Additional components include, for example, polyvalent metal cations, suspending agents, and other additional components.

Polyvalent Metal Cations

Suitable polyvalent metal cations include divalent and trivalent metals, divalent metals being preferred. Exemplary metal cations include alkaline earth metals, such as magnesium, calcium, zinc, and copper, and trivalent metals such as aluminum and iron. Preferred are calcium and magnesium.

The polyvalent metal cation can be added as an inorganic salt, organic salt, or as a hydroxide. The polyvalent metal cation may also be added as a salt with anionic surfactant, including the primary anionic surfactants as mentioned above, or detergents as mentioned below.

Preferably, the polyvalent metal cation is introduced as an inorganic salt or organic salt. Inorganic salts include chloride, bromide, iodine, nitrate, or sulfate, more preferably chloride or sulfate. Organic salts include L-glutamate, lactate, malate, succinate, acetate, fumarate, L-glutamic acid hydrochloride, and tartarate.

It will be clear to those skilled in the art that, if polyvalent salts of the anionic surfactant is used as the mode of introducing the polyvalent metal cations into the compositions hereof, only a fraction of the anionic surfactant may be of polyvalent form, the remainder of the anionic surfactant being necessarily added in monovalent form.

Hardness of the conditioning shampoo compositions can be measured by standard methods in the art, such as by ethylene diamine tetraacetic acid (EDTA) titration. In the event that the composition contains dyes or other color materials that interfere with the ability of EDTA titration to yield a perceptible color change, hardness should be determined from the composition in the absence of the interfering dye or color.

Suspending Agents

A preferred additional component is a suspending agent, particularly for compositions comprising silicone compounds of high viscosity and/or large particle size. When present, the suspending agent is in dispersed form in the compositions. The suspending agent will generally comprise from about 0.1% to about 10%, and more typically from about 0.3% to about 5.0%, by weight, of the composition.

Preferred suspending agents include acyl derivatives such as ethylene glycol stearates, both mono and distearate, long chain amine oxides such as alkyl (C16-C22) dimethyl amine oxides, e.g., stearyl dimethyl amine oxide, and mixtures thereof. When used in the shampoo compositions, these preferred suspending agents are present in the composition in crystalline form. These suspending agents are described in U.S. Patent 4,741,855.

Other suitable suspending agents include alkanol amides of fatty acids, preferably having from about 16 to about 22 carbon atoms, more preferably about 16 to 18 carbon atoms, preferred examples of which include stearic monoethanolamide, cocomonoethanolamide, stearic diethanolamide, stearic monoisopropanolamide and stearic monoethanolamide stearate.

Other suitable suspending agents include N,N-dihydrocarbyl amido benzoic acid and soluble salts thereof (e.g., Na and K salts), particularly N,N-di(hydrogenated) C16, C18 and tallow amido benzoic acid species of this family, which are commercially available from Stepan Company (Northfield, Illinois, USA).

Other suitable suspending agents include xanthan gum. The use of xanthan gum as a suspending agent in silicone containing shampoo compositions is described, for example, in U.S. Patent 4,788,006, which is incorporated herein by reference in its entirety. Combinations of long chain acyl derivatives and xanthan gum may also be used as a suspending agent in the shampoo compositions. Such combinations are described in U.S. Patent 4,704,272, which is incorporated herein by reference in its entirety.

Other suitable suspending agents include carboxyvinyl polymers. Preferred among these polymers are the copolymers of acrylic acid crosslinked with polyallylsucrose as described in U.S. Patent 2,798,053, which is incorporated herein by reference in its entirety. Examples of these polymers include the carbomers, which are homopolymers of acrylic acid crosslinked with an allyl ether of pentaerythrotol, an allyl ether of sucrose, or an allyl ether of

propylene. Neutralizers may be required, for example, amino methyl propanol, triethanol amine, or sodium hydroxide.

Other suitable suspending agents can be used in the compositions, including those that can impart a gel-like viscosity to the composition, such as water soluble or colloidally water soluble polymers like cellulose ethers such as hydroxyethyl cellulose, hydroxymethyl cellulose, hydroxypropyl cellulose, and materials such as guar gum, polyvinyl alcohol, polyvinyl pyrrolidone, hydroxypropyl guar gum, starch and starch derivatives.

Other Additional Components

A wide variety of other additional ingredients can be formulated into the present compositions. These include: other conditioning agents such as hydrolysed collagen with tradename Peptein 2000 available from Hormel, vitamin E with tradename Emix-d available from Eisai, panthenol available from Roche, panthenyl ethyl ether available from Roche, hydrolysed keratin, proteins, plant extracts, and nutrients; hair-fixative polymers such as amphoteric fixative polymers, cationic fixative polymers, anionic fixative polymers, nonionic fixative polymers, and silicone grafted copolymers; preservatives such as benzyl alcohol, methyl paraben, propyl paraben and imidazolidinyl urea; pH adjusting agents, such as citric acid, sodium citrate, succinic acid, phosphoric acid, sodium hydroxide, sodium carbonate; salts, in general, such as potassium acetate and sodium chloride; coloring agents, such as any of the FD&C or D&C dyes; hair oxidizing (bleaching) agents, such as hydrogen peroxide, perborate and persulfate salts; hair reducing agents such as the thioglycolates; perfumes; and sequestering agents, such as disodium ethylenediamine tetra-acetate; ultraviolet and infrared screening and absorbing agents such as octyl salicylate, and antidandruff agents such zinc pyridinethione. Such optional ingredients generally are used individually at levels from about 0.001% to about 10.0%, preferably from about 0.01% to about 5.0% by weight of the composition.

EXAMPLES

The following examples further describe and demonstrate embodiments within the scope of the present invention. The examples are given solely for the purpose of illustration and are not to be construed as limitations of the present invention, as many variations thereof are possible without departing from the spirit and scope of the invention. Ingredients are identified by chemical or CTFA name, or otherwise defined below.

Method of Preparation

The hair spray, mousses, lotions and conditioners are suitably made as follows: If included in the formula, polymeric materials are dispersed in water at room temperature. If required, the polymeric materials such as Carbomer and Acrylates/Stearth-20 Methacrylate Copolymer may be neutralized after dispersing. The mixture is then heated up to above 60°C, and fatty alcohols and emulsifiers are added if included in the formulation. After cooling down to below 50°C, the remaining components are added with agitation then cooled down to about 30°C. Ethanol is added here if included in the formula. A triblender and mill can be used if necessary to disperse the materials. As appropriate, the mixture thus obtained can be packed into an aerosol can with propellant.

The shampoos are suitably prepared by any conventional method well known in the art. A suitable method is as follows: polymer and surfactants are dispersed in water to form a homogeneous mixture. To this mixture are added the other ingredients except for silicone emulsion (if present), perfume, and salt; the obtained mixture is agitated. If present, the silicone emulsion is made with Dimethicone or Dimethiconol, a small amount of detergentsurfactant, and a portion of water. The obtained mixture is then passed through a heat exchanger to cool, and the silicone emulsion, perfume, and salt are added. The obtained compositions are poured into bottles to make hair shampoo compositions.

Alternatively, water and surfactants and any other solids that need to be melted can be mixed together at elevated temperature, e.g., above about 70°C, to speed the mixing into shampoo. Additional ingredients can be added either to this hot premix or after cooling the premix. The ingredients are mixed thoroughly at the elevated temperature and then pumped through a high shear mill and then through a heat exchanger to cool them to ambient temperature. If present in the composition, silicone emulsified at room temperature in concentrated surfactant is added to the cooled mix.

Compositions

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Ex. 6
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	1.0	-	-	-	-	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)-amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	0.8	-	-	-	-
4-methyl-6,7-dihydroxycoumarin *3	-	-	1.0	-	-	-
4-methyl-7,7'-dimethylamino coumarin *4	-	-	-	0.1	-	-

Hydrolyzed Collagen *40	0.01	-	-	-	-	-
Vitamin E *41	0.01	-	-	-	-	-
Panthenol *42	0.025	-	-	-	-	-
Panthenyl Ethyl Ether *43	0.225	-	-	-	-	-
Deionized Water	q.s. to 100%					

	Ex. 7	Ex. 8	Ex. 9	Ex. 10
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.8	1.0	0.5	0.8
Silicone Dow Corning 200	2.0	-	-	-
Silicone Emulsion *15	-	2.0	2.0	1.0
Ammonium Laureth-3 Sulfate	12.0	12.0	15.0	10.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0
Sodium Lauroyl Sarcosinate *10	-	-	-	1.0
Cocamidopropylbetaine *11	-	-	-	2.0
Cocamide MEA	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Deionized Water	q.s. to 100%			

	Ex.11	Ex.12	Ex.13	Ex.14	Ex.15
Disodium-1,4-bis(2-sulfostyryl)biphenyl *1	0.80	-	-	0.10	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino 1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	-	-	-	0.80
1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline *5	-	-	-	0.60	-
4,4'-bis(5-methylbenzoxazol-2-yl)stilbene *25	-	-	0.50	-	-
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	-	0.20	-	-	-
Silicone Emulsion *15	1.00	0.50	2.00	0.20	1.00
Cetyl Alcohol *19	2.00	0.50	0.60	0.20	0.60
Stearyl Alcohol *20	3.00	0.50	0.40	0.20	0.40
Behenyl Alcohol *21	-	-	-	0.20	-

Polysorbate 60 *26	1.00	-	-	-	-
Lauryl Methyl Gluceth-10 Hydroxypropyl-dimonium Chloride *27	-	1.00	-	0.50	-
Dihydrogenated Tallowamidoethyl Hydroxyethylmonium Methosulfate *28	-	-	0.10	-	0.10
Citric Acid *29	0-0.2	0-0.2	0-0.2	0-0.2	-
Hydroxyl Propyl Cellulose *30	-	-	-	-	0.50
Polyoxyethylene Glycol *16	0.50	0.50	0.50	0.50	0.50
Aminomethylpropanol *31	2.10	-	2.10	-	-
Acrylates/Steareth-20 Methacrylate Copolymer *32	1.75	-	1.75	-	-
Cetyl Hydroxyethyl Cellulose *33	-	0.75	-	0.75	-
Polyquaternium-10 *18	0.10	-	0.10	0.20	-
Polyquaternium-7 *34	-	0.50	-	-	1.00
Pentaerythritol Tetraisostearate *22	1.00	3.00	-	-	-
Trimethylolpropane Trioleate *24	-	-	0.50	-	0.10
Trimethylolpropane Triisostearate *23	-	-	-	0.20	-
Preservatives	0.90	0.90	0.90	0.90	-
Perfume	0.08	0.08	0.08	0.08	0.08
Glycerin *7	-	-	5.00	-	-
Propylene Glycol *8	-	-	-	5.00	-
Denatured Ethyl Alcohol	-	-	-	-	18.98
Deionized Water	q.s. to 100%				

	Ex.16	Ex.17	Ex.18	Ex.19
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	0.80	-	-	-
4-methyl-6,7-dihydroxycoumarin *3	-	0.50	-	-
4-methyl-7,7'-dimethylamino coumarin *4	-	-	0.50	-
2-(4-styryl-3-sulfophenyl)-2H-naphto[1,2-d]triazole	-	-	-	0.40
15/85 Silicone Blend *35	-	-	-	4.20
Silicone Emulsion *15	0.20	0.50	1.00	-
Cetyl Alcohol *19	0.20	-	2.00	2.50

Stearyl Alcohol *20	0.20	-	-	4.50
Behenyl Alcohol *21	0.10	0.50	-	-
Stearamidopropyl dimethylamine *36	0.20	-	-	2.00
L-Glutamic Acid *37	0.08	-	-	0.64
Hydroxyethyl Cellulose *17	-	0.50	-	-
Polyoxyethylene Glycol *16	0.50	0.50	0.50	0.20
Carbomer *38	0.50	-	0.50	-
Aminomethylpropanol *31	0.60	-	0.60	-
Polyquaternium-10 *18	1.00	0.10	-	0.20
Polyquaternium-7 *34	-	-	0.50	-
Pentaerythritol Tetraisostearate *22	0.10	0.10	0.20	-
Trimethylolpropane Trioleate *24	0.10	0.10	0.05	-
Trimethylolpropane Triisostearate *23	-	-	-	0.25
Preservatives	-	-	-	0.53
Perfume	0.08	0.08	0.08	0.20
Glycerin *7	-	5.00	-	-
Propylene Glycol *8	-	-	5.00	-
Denatured Ethyl Alcohol	19.07	18.42	17.81	-
Deionized Water	q.s. to 100%			

	Ex. 20	Ex. 21	Ex. 22	Ex. 23	Ex. 24	Ex. 25
Disodium-1,4-bis(2-sulfostyryl)biphenyl *1	0.80	1.60	0.20	1.60	0.20	0.20
15/85 Silicone Blend *35	-	-	-	-	4.20	4.20
Silicone Emulsion *15	1.00	1.00	1.00	0.50	-	-
Cetyl Alcohol *19	-	-	-	-	2.50	2.50
Stearyl Alcohol *20	-	-	-	-	4.50	4.50
Alkyl Trimethyl Ammonium Chloride *39	-	-	-	-	-	2.50
Stearamidopropyl dimethylamine *36	-	-	-	-	2.00	-
L-Glutamic Acid *37	-	-	-	-	0.64	-
Preservatives	0.90	0.90	-	-	0.53	0.53
Perfume	0.08	0.08	0.08	0.08	0.20	0.20
Denatured Ethyl Alcohol	-	-	19.74	19.56	-	-

Deionized Water	-----q.s. to 100%-----
-----------------	------------------------

Components	Ex.26	Ex.27	Ex.28
4-methyl-7,7'-dimethylamino coumarin *4	0.1	-	-
2-(4-styryl-3-sulfophenyl)-2H-naphto[1,2-d]triazole	1.0	-	-
1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline *5	-	0.5	-
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	-	-	1.0
Polyquaternium-10 *18	-	0.35	0.35
Monoalkyl trimethyl ammonium	0.2	-	-
Ammonium Laureth-3 Sulfate	15.0	10.0	10.0
Ammonium Lauryl Sulfate	2.0	-	-
N-acyl-L-gultamate Triethanolamine *9	-	2.0	2.0
Sodium Lauroyl Sarcosinate *10	2.0	2.0	1.0
Cocamidopropylbetaine *11	4.0	2.0	2.0
Laureth-20 *12	0.5	0.5	-
Ethylene Glycol Distearate *14	2.0	2.0	1.5
Cocamide MEA	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2
Propylene Glycol *8	10.0	5.0	10.0
Silicone	3.0	1.6	2.0
Polyoxyethylene Glycol *16	0.1	0.5	-
Hydroxyethyl Cellulose *17	-	-	0.5
Cetyl alcohol *19	0.1	0.1	-
Stearyl alcohol *20	0.1	0.2	-
Perfume	0.5	0.5	0.5
Pentaerythritol Tetraisostearate *22	0.5	-	-
Trimethylolpropane Triisostearate *23	-	0.3	-
Trimethylolpropane Trioleate *24	-	-	0.5
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5
Deionized Water	-----q.s. to 100%-----		

Components	Ex. 29	Ex. 30	Ex. 31	Ex. 32
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.2	0.8	0.5	0.3

Polyquaternium-10 *18	0.4	0.5	-	-
Monoalkyl trimethyl ammonium	-	-	0.1	0.2
Ammonium Laureth-3 Sulfate	12.0	12.0	12.0	12.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0
Cocamide MEA	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Deionized Water	-----q.s. to 100%-----			

Components	Ex. 33	Ex. 34	Ex. 35	Ex. 36	Ex. 37	Ex. 38
Disodium-1,4-bis(2-sulfostyryl)biphenyl *1	0.20	0.80	0.20	-	0.20	0.80
4-methyl-6,7-dihydroxycoumarin *3	-	-	-	0.80	-	-
1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline *5	-	-	0.40	-	-	-
Lauryl Methyl Gluceth-10	-	1.00	-	-	-	-
Hydroxypropyl-dimonium Chloride *27						
Alkyl Trimethyl Ammonium Chloride *39	-	-	-	1.00	-	-
Dihydrogenated Tallowamidoethyl Hydroxyethylmonium Methosulfate *28	-	-	0.25	2.00	-	0.20
Stearamidopropyl dimethylamine *36	2.00	2.00	2.00	-	-	-
L-Glutamic Acid *37	0.64	0.64	0.80	-	-	-
Polyquaternium-10 *18	0.40	-	-	0.40	0.40	0.40
Polyquaternium-7 *34	-	0.40	-	-	-	-
Preservatives	0.53	0.53	0.53	0.53	-	-
Perfume	0.08	0.08	0.08	0.08	0.08	0.08
Denatured Ethyl Alcohol	-	-	-	-	19.86	19.70
Deionized Water	-----q.s. to 100%-----					

	Ex.39
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	1.0
Staryl alcohol *20	0.2
Ammonium Laureth-3 Sulfate	10.0
N-acyl-L-gultamate Triethanolamine *9	2.0
Sodium Lauroyl Sarcosinate *10	1.0
Cocamidopropylbetaine *11	2.0
Ethylene Glycol Distearate *14	1.5
Cocamide MEA	1.5
DMDM Hydantoin	0.2
Propylene Glycol *8	10.0
Silicone	2.0
Hydroxyethyl Cellulose *17	0.5
Perfume	0.5
Trimethylolpropane Trioleate *24	0.5
Monosodium phosphate	0 to 0.5
Disodium phosphate	0 to 0.5
Deionized Water	q.s. to 100%

	Ex. 40	Ex. 41	Ex. 42	Ex. 43
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.8	1.0	0.5	0.8
Cetyl alcohol *19	0.1	0.1	-	-
Staryl alcohol *20	0.1	-	0.1	-
Behenyl alcohol *21	-	-	-	0.2
Ammonium Laureth-3 Sulfate	12.0	12.0	15.0	10.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0
Sodium Lauroyl Sarcosinate	-	-	-	1.0
Cocamidopropylbetaine	-	-	-	2.0
Cocamide MEA	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Deionized Water	q.s to 100%			

	Ex. 44	Ex. 45	Ex. 46	Ex. 47
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.20	0.50	0.80	0.50
Cetyl Alcohol *19	0.40	-	1.00	0.50
Stearyl Alcohol *20	0.60	1.00	-	0.50
Polysorbate 60 *26	0.20	0.20	0.20	-
Preservatives	0.53	0.53	0.53	-
Perfume	0.08	0.08	0.08	0.08
Denatured Ethyl Alcohol	-	-	-	19.68
Deionized Water	-----q.s. to 100%-----			

	Ex. 48	Ex. 49	Ex. 50
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.80	-	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	0.50	1.00
Cetyl Alcohol *19	2.00	2.50	2.50
Stearyl Alcohol *20	-	4.50	4.50
15/85 Silicone Blend *35	-	4.20	4.20
Polysorbate 60 *26	-	-	-
Alkyl Trimethyl Ammonium Chloride *39	-	-	2.50
Stearamidopropyl dimethylamine *36	-	2.00	-
L-Glutamic Acid *37	-	0.64	-
Preservatives	-	0.53	0.53
Perfume	0.08	0.20	0.20
Denatured Ethyl Alcohol	19.42	-	-
Deionized Water	-----q.s. to 100%-----		

	EX. 51	EX. 52	EX. 53	EX. 54	EX. 55
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	1.0	-	-	-	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)-amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	0.8	-	-	-

4-methyl-6,7-dihydroxycoumarin *3	-	-	1.0	-	-
4-methyl-7,7'-dimethylamino coumarin *4	-	-	-	0.1	-
2-(4-styryl-3-sulfophenyl)-2H-naphto[1,2-d]triazole	-	-	-	1.0	-
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	-	-	-	-	1.0
Citronellol	0.2	-	-	-	0.05
α -ionone	-	0.2	-	-	-
Benzaldehyde	0.05	0.05	0.2	-	-
Geraniol	-	-	0.05	0.2	-
Eugenol	-	-	-	0.05	0.2
Ammonium Laureth-3 Sulfate	12	12	12	12	12
Ammonium Lauryl Sulfate	2	2	2	2	2
N-acyl-L-gultamate	2	2	2	2	2
Triethanolamine *9					
Cocamidopropylbetaine *11	2	2	2	2	2
Ethylene Glycol Distearate *14	2.0	2.0	2.0	2.0	2.0
Cocamide MEA	1.5	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2	0.2
MgCl ₂ .6H ₂ O	0.5	0.5	0.5	0.5	0.5
Monosodium phosphate	0-1.0	0-1.0	0-1.0	0-1.0	0-1.0
Disodium phosphate	0-1.0	0-1.0	0-1.0	0-1.0	0-1.0
Polyquaternium-10 *18	0.5	0.5	0.5	0.5	0.5
Silicone	2.0	2.0	2.0	2.0	2.0
Deionized Water	----- q.s. to 100% -----				
pH	4.0-6.5	4.0-6.5	4.0-6.5	4.0-6.5	4.0-6.5
Hydrolyzed Collagen *40	0.01	-	-	-	-
Vitamin E *41	0.01	-	-	-	-
Panthenol *42	0.025	-	-	-	-
Panthenyl Ethyl Ether *43	0.225	-	-	-	-

	Ex. 56	Ex. 57	Ex. 58	Ex. 59	Ex. 60
--	--------	--------	--------	--------	--------

Disodium-1,4-bis(2-sulfostyryl) biphenyl*1	-	1.50	-	-	1.50
6,7-dihydroxy Coumarin	-	-	0.20	-	-
2-(4-styryl-3-sulfophenyl)-2-naphtotriazole	-	-	-	0.40	-
1,3,-Diphenyl-pyrazoline	0.50	-	-	-	-
Citronellol	0.08	-	0.20	-	-
α -ionone	-	0.08	-	0.05	-
Benzaldehyde	-	-	0.10	-	1.00
Geraniol	-	-	-	0.2	-
Eugenol	-	-	-	-	0.1
Cetyl Alcohol*19	2.00	-	-	2.50	-
Stearyl Alcohol*20	3.00	-	-	4.50	-
Behenyl Alcohol*21	-	-	0.50	-	-
15/85 Silicone Blend*35	-	-	-	4.20	-
Silicone Emulsion*15	1.00	1.00	0.50	-	-
Polysorbate 60*26	1.00	-	-	-	-
Stearamidopropyl dimethylamine*36	-	-	-	2.00	-
L-Glutamic Acid*37	-	-	-	0.64	-
Citric Acid*29	0-0.2	0-0.2	-	-	-
Hydroxyethyl Cellulose*17	-	-	0.50	-	-
Polyoxyethylene Glycol*16	0.50	-	0.50	0.20	-
Aminomethylpropanol*31	2.10	-	-	-	-
Acrylates/Steareth-20 Methacrylate Copolymer*32	1.75	-	-	-	-
Cetyl Hydroxyethyl Cellulose*33	-	0.80	-	-	-
Polyquaternium-10*18	0.10	-	0.10	0.20	-
Pentaerythritol Tetraisostearate*22	1.00	-	-	-	-
Preservatives	0.90	0.53	0.90	0.53	0.53
Glycerin*7	-	-	5.00	-	-
Propylene Glycol*8	-	5.00	-	-	-
Denatured Ethyl Alcohol	-	-	18.34	-	-
Deionized Water	q.s. to 100%				

	Ex. 61	Ex. 62	Ex. 63	Ex. 64
--	--------	--------	--------	--------

Disodium-1,4-bis(2-sulfostyryl)biphenyl*1	0.8	1.0	0.5	0.8
Pentaerythritol Tetraisostearate*22	0.3	-	-	-
Trimethylolpropane Triisostearate*23	-	0.5	-	-
Pentaerythritol Tetraoleate	-	-	0.3	-
Trimethylolpropane Trioleate*24	-	-	-	0.8
Ammonium Laureth-3 Sulfate	12.0	12.0	15.0	10.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0
Sodium Lauroyl Sarcosinate*10	-	-	-	1.0
Cocamidopropylbetaine*11	-	-	-	2.0
Cocamide MEA	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Deionized Water	-----q.s to 100%-----			

	Ex. 65	Ex. 66	Ex. 67	Ex. 68
Disodium-1,4-bis(2-sulfostyryl)biphenyl*1	0.80	0.80	1.60	0.20
Pentaerythritol Tetraisostearate *22	1.00	0.50	-	-
Trimethylolpropane Trioleate *24	-	-	0.50	-
Trimethylolpropane Triisostearate *23	-	-	-	0.50
Preservatives	0.53	0.53	0.53	0.53
Perfume	0.08	0.08	0.08	0.08
Deionized Water	-----q.s. to 100%-----			

Definitions

- *1 Disodium-1,4-bis(2-sulfostyryl)biphenyl: TINOPAL CBX obtained by Ciba Geigy.
- *2 4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids: Tinopal UNPA-GX obtained by Ciba Geigy.
- *3 4-methyl-6,7-dihydroxycoumarin: available from Wako.
- *4 4-methyl-7,7'-dimethylamino coumarin: available from Wako.
- *5 1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline: Blankophor DCB obtained by Bayer.
- *6 2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine: available from Ciba Geigy.

- *7 Glycerin: Glycerin USP obtained by Nihon Yushi.
- *8 Propylene Glycol: Lexol PG-865 obtained by Inolex.
- *9 N-acyl-L-gultamate Triethanolamine: CT12S obtained by Ajinomoto.
- *10 Sodium Lauroyl Sarcosinate: Soypon obtained by Kawaken Fine Chem.
- *11 Cocamidopropylbetaine: Tegobetain obtained by Th.Goldschmidt AG.
- *12 Laureth-20: BL-20 obtained by Nikko.
- *13 Alkyl polyglucoside: Plantacare 2000UP obtained by Henkel.
- *14 Ethylene Glycol Distearate: EGDS obtained by Th. Goldschmidt AG.
- *15 Silicone Emulsion: X65-4829 obtained by Tosil/GE.
- *16 Polyoxyethylene Glycol: WSR N-10 obtained by Amerchol.
- *17 Hydroxyethyl Cellulose: available from Aqualon.
- *18 Polyquaternium-10: UCARE Polymer LR 400 obtained by Amerchol.
- *19 Cetyl Alcohol: Konol series obtained by Shinihon Rika.
- *20 Stearyl Alcohol: Konol series obtained by Shinihon Rika.
- *21 Behenyl Alcohol: 1-Docosanol (97%) obtained by Wako.
- *22 Pentaerythritol Tetraisostearate: KAK PTI obtained by Kokyu alcohol.
- *23 Trimethylolpropane Triisostearate: KAK TTI obtained by Kokyu alcohol.
- *24 Trimethylolpropane Trioleate: Enujerubu TP3SO obtained by Shinihon Rika.
- *25 4,4'-bis(5-methylbenzoxazol-2-yl)stilbene: available from TCI.
- *26 Polysorbate 60: Tween 60 obtained by ICI.
- *27 Lauryl Methyl Gluceth-10 Hydroxypropylmonium Chloride: Glucquat 125 obtained by Amerchol.
- *28 Dihydrogenated Tallowamidoethyl Hydroxyethylmonium Methosulfate: Varisoft 110 obtained by Witco.
- *29 Citric Acid: Anhydrous Citric acid obtained by Haarman & Reimer.
- *30 Hydroxyl Propyl Cellulose: Metlose obtained by Shinetsu.
- *31 Aminomethylpropanol: AMP-regular obtained by Angus.
- *32 Acrylates/Steareth-20 Methacrylate Copolymer: Aculyn obtained by Rohm&Haas.
- *33 Cetyl Hydroxyethyl Cellulose: Polysurf 67 obtained by Aqualon.
- *34 Polyquaternium-7: Merquat S obtained by Calgon.
- *35 15/85 Silicone Blend: available from Shinetsu.
- *36 Stearamidopropyl dimethylamine: Amidoamine MPS obtained by Nikko.
- *37 L-Glutamic Acid: L-Glutamic acid (cosmetic grade) obtained by Ajinomoto.

- *38 Carbomer: Carbopol 980 obtained by BF Goodrich.
- *39 Alkyl Trimethyl Ammonium Chloride: available from TOHO.
- *40 Hydrolyzed Collagen: PEPTEIN 2000 obtained by Hormel.
- *41 Vitamin E: EMIX-d obtained by Eisai.
- *42 Panthenol: available from Roche.
- *43 Panthenyl Ethyl Ether: available from Roche.

The embodiments disclosed and represented by the previous examples have many advantages. For example, they can provide the benefits associated with optical brighteners (e.g., shine, UV protection, color alteration) while also providing desirable conditioning benefits.

It is understood that the examples and embodiments described herein are for illustrative purposes only and that various modifications or changes in light thereof will be suggested to one skilled in the art without departing from its spirit and scope.

What is claimed is:

1. A hair care composition comprising:
 - (a) an effective amount of an optical brightener; and
 - (b) a hair conditioning agent selected from the group consisting of silicone compounds, cationic compounds, high melting point compounds, perfume compounds, water-insoluble high molecular weight oily compounds, and mixtures thereof.
2. A hair care composition comprising:
 - (a) an effective amount of an optical brightener selected from the group consisting of polystyrylstilbenes, hydroxycoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof; and
 - (b) a silicone compound.
3. The hair care composition of claim 2 further comprising a carrier suitable for application to the hair.
4. The hair care composition of claim 2 wherein the optical brightener is a polystyrylstilbene.
5. The hair care composition of claim 2 wherein the optical brightener is a hydroxycoumarin.
6. A hair care composition comprising:
 - (a) an effective amount of an optical brightener; and
 - (b) a silicone compound selected from the group consisting of high molecular weight dimethicones.
7. The hair care composition of claim 6 further comprising a carrier suitable for application to the hair.
8. The hair care composition of claim 7 wherein the optical brightener is a polystyrylstilbene.

9. The hair care composition of claim 7 wherein the optical brightener is a hydroxycoumarin.
10. The hair care composition of claim 7 wherein the optical brightener is a triazinstilbene.
11. The hair care composition of claim 2 or claim 6 containing from about 0.001 wt% to about 20 wt% of the optical brightener.
12. The hair care composition of claim 11 containing from about 0.01 wt% to about 15 wt% of the silicone compound.
13. The hair care composition of any of claims 2 - 12 further comprising another conditioning agent.
14. A shampoo composition comprising:
 - (a) an effective amount of an optical brightener;
 - (b) a deterotive surfactant suitable for cleansing the hair; and
 - (c) a silicone compound.
15. The shampoo composition of claim 14 further comprising a carrier suitable for application to the hair.
16. The shampoo composition of claim 15 wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof.
17. The shampoo composition of claim 16 containing from about 0.001 wt% to about 20 wt% of the optical brightener.
18. The shampoo composition of claim 17 containing from about 0.01 wt% to about 15 wt% of the silicone compound.
19. The shampoo composition of any of claims 2 - 18 further comprising another conditioning agent.

20. A hair care composition comprising:
 - (a) an effective amount of an optical brightener;
 - (b) a cationic compound;
 - (c) a carrier suitable for application to hair.
21. The composition of claim 20 further comprising a deterotive surfactant suitable for cleansing hair.
22. The composition of claim 21 further comprising another conditioning agent.
23. The composition of claim 20 further comprising another conditioning agent.
24. The composition of any of claims 20 - 23 wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof.
25. The composition of claim 24 containing from about 0.001 wt% to about 20 wt% of the optical brightener.
26. The composition of claim 25 containing from about 0.01 wt% to about 10 wt% of the cationic compound.
27. A hair care composition comprising:
 - (a) an effective amount of an optical brightener;
 - (b) a high melting point compound; and
 - (c) a carrier suitable for application to the hair;
28. The hair care composition of claim 27 further comprising a deterotive surfactant suitable for cleansing the hair.
29. The hair care composition of claim 28 further comprising another conditioning agent.

30. The composition of claim 27 further comprising another conditioning agent.
31. The hair care composition of any claims 27 - 30 wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof.
32. The hair care composition of claim 31 containing from about 0.001 wt% to about 20 wt% of the optical brightener.
33. The composition of claim 32 containing from about 0.01 wt% to about 20 wt% of the high melting point compound.
34. The composition of claim 33 wherein the high melting point compound is selected from the group consisting of cetyl alcohol, stearyl alcohol, behenyl alcohol, oleyl alcohol, and mixtures thereof.
35. A hair care composition comprising:
 - (a) an effective amount of an optical brightener;
 - (b) perfume compound having a CLogP of between about -1.0 and about 8.0; and
 - (c) a carrier suitable for application to the hair.
36. The composition of claim 35 further comprising a deterotive surfactant suitable for cleansing the hair.
37. The composition of claim 36 further comprising another conditioning agent.
38. The composition of claim 35 further comprising another conditioning agent.

39. The composition of any of claims 35 - 38 wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof.

40. The composition of claim 39 containing from about 0.001 wt% to about 20 wt% of the optical brightener.

41. The composition of claim 40 containing at least about 0.05% of the perfume compound.

42. The composition of claim 41 wherein the perfume compound is selected from the group consisting of a carbonyl containing perfume compound, an ether containing perfume compound, a hydroxy containing perfume compound, and mixtures thereof.

43. A hair care composition comprising:

- (a) an effective amount of an optical brightener; and
- (b) a water-insoluble high molecular weight oily compound.

44. The hair care composition of claim 43 further comprising a carrier suitable for application to hair.

45. The composition of claim 44 further comprising a deterersive surfactant suitable for cleansing the hair.

46. The composition of claim 45 further comprising another conditioning agent.

47. The composition of any of claims 43 - 46 wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, and imidazoles, mixtures thereof.

48. The composition of claim 48 containing from about 0.001 wt% to about 20 wt% of the optical brightener.

49. The composition of claim 48 containing from about 0.1 wt% to about 10 wt% of the water-insoluble high molecular weight oily compound.

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : A61K 7/06, 7/13</p>		<p>A3</p>	<p>(11) International Publication Number: WO 99/13823 (43) International Publication Date: 25 March 1999 (25.03.99)</p>
<p>(21) International Application Number: PCT/IB98/01381 (22) International Filing Date: 4 September 1998 (04.09.98)</p>		<p>(74) Agents: REED, T., David et al.; The Procter & Gamble Company, 5299 Spring Grove Avenue, Cincinnati, OH 45217-1087 (US).</p>	
<p>(30) Priority Data: PCT/US97/16412 17 September 1997 (17.09.97) US PCT/US97/16617 17 September 1997 (17.09.97) US PCT/US97/16618 17 September 1997 (17.09.97) US PCT/US97/16619 17 September 1997 (17.09.97) US PCT/US97/16615 17 September 1997 (17.09.97) US</p>		<p>(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p>	
<p>(71) Applicant (for all designated States except US): THE PROCTER & GAMBLE COMPANY [US/US]; One Procter & Gamble Plaza, Cincinnati, OH 45202 (US).</p>		<p>Published With international search report.</p>	
<p>(72) Inventors; and (75) Inventors/Applicants (for US only): MITSUMATSU, Arata [JP/JP]; 6-16-306, Midorigaoka-cho, Ashiya-shi, Hyogo 659 (JP). LUO, Xiaochun [JP/JP]; 5-15-703, Koyo-cho Naka, Higashinada-ku, Kobe 658 (JP). UCHIYAMA, Hirotaka [JP/JP]; 5-8-17-3B, Nishi-Okamoto, Higashinada-ku, Kobe 658 (JP). SETSER, Julia, Leet [US/JP]; 2101, 5-15, Koyo-cho Naka, Higashinada-ku, Kobe 658 (JP).</p>		<p>(88) Date of publication of the international search report: 12 August 1999 (12.08.99)</p>	
<p>(54) Title: HAIR CARE COMPOSITIONS COMPRISING OPTICAL BRIGHTENERS AND HAIR CONDITIONING AGENTS</p>			
<p>(57) Abstract</p> <p>Disclosed are hair care compositions comprising: (a) an effective amount of an optical brightener; and (b) a hair conditioning agent selected from the group consisting of silicone compounds, cationic compounds, high melting point compounds, perfume compounds, water-insoluble high molecular weight oily compounds, and mixtures thereof.</p>			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

INTERNATIONAL SEARCH REPORT

International Application No
PCT/IB 98/01381

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61K7/06 A61K7/13

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 94 06409 A (COLGATE PALMOLIVE CO) 31 March 1994 see examples 1,2 ---	1-3,6,7, 11,13
X	EP 0 717 978 A (CURTIS HELENE IND INC) 26 June 1996 see page 21, line 1-50; claims 1,13,35; examples 35,72 ---	1-4,6-8, 10-13
X	GB 2 307 639 A (GEN ELECTRIC) 4 June 1997 cited in the application see page 8, line 3 - page 9, line 14; claims ---	1,6,7, 11-13
P,X	EP 0 834 303 A (KAO CORP) 8 April 1998 see examples 9,13 ---	1,6,7, 11-13 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

1 February 1999

Date of mailing of the international search report

06.05.99

Name and mailing address of the ISA
European Patent Office, P.B. 5818 Patenttaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl
Fax: (+31-70) 340-3016

Authorized officer

Veronese, A

INTERNATIONAL SEARCH REPORT

International Application No
PCT/IB 98/01381

C(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 733 355 A (KAO CORP) 25 September 1996 see example 2 ---	1,6,7, 11-13
X	GB 1 328 108 A (UNILEVER LTD) 30 August 1973 cited in the application see page 1, line 65-92; claim 6 ---	1,6,7, 11,12
X	US 3 658 985 A (OLSON FRANK WESLEY JR ET AL) 25 April 1972 cited in the application see page 1, column 1, line 63 - page 1, column 2, line 27; examples 3-5 ---	1-3,11, 12
X	CA 1 255 603 A (COSMEPRO INC LAB) 13 June 1989 cited in the application see page 3, line 13 - page 4, line 9; claim 12 ---	1
X	US 4 312 855 A (GRAND PAUL S) 26 January 1982 cited in the application see column 5, line 45 - column 7, line 2 see claims 1-4,9; example 11 * see column 17: Table III *	1
A	EP 0 336 709 A (DOW CORNING) 11 October 1989 see claims 2,5,20-22 ---	2
A	DE 26 32 810 A (HENKEL KGAA) 26 January 1978 see examples -----	2

INTERNATIONAL SEARCH REPORT

International application No.

PCT/IB 98/01381

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1 partial, 2-13 complete

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Subject	claims	Description
1	1 part 2-13 complete	A hair care composition comprising a) an effective amount of an optical brightener. b) a silicon compound
2	1 part 14-19 complete	As far as not comprised in invention 1 a) an effective amount of an optical brightener. b) a deterutive surfactant. c) a silicon compound
3	1 part 20-26 complete	As far as not comprised in inv 1-2 a) an effective amount of an optical brightener. b) a cationic compound.
4	1 part 27-34 complete	As far as not comprised in inv 1-3 A hair care composition comprising a) an effective amount of an optical brightener. b) a high melting point compound.
5	1 part 35-42 compl	As far as not comprised in inv 1-4 A hair care composition comprising a) an effective amount of an optical brightener. b) a perfume compound.
6	1 part 43-49 compl	As far as not comprised in inv 1-5 a) an effective amount of an optical brightener. b) a water-insoluble high molecular weight oily compound

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/IB 98/01381

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9406409	A	31-03-1994	AU 4928393 A		12-04-1994
			CN 1089830 A		27-07-1994
			MX 9305746 A		31-05-1994
			US 5580494 A		03-12-1996
			ZA 9306928 A		20-03-1995
EP 0717978	A	26-06-1996	US 5589177 A		31-12-1996
			AU 4026395 A		13-06-1996
			AU 688632 B		12-03-1998
			AU 4027195 A		13-06-1996
			CA 2163854 A		07-06-1996
			CN 1138982 A		01-01-1997
			FI 955866 A		07-06-1996
			NO 954916 A		07-06-1996
			NZ 280599 A		27-07-1997
			ZA 9510058 A		04-06-1996
GB 2307639	A	04-06-1997	DE 19646804 A		22-05-1997
			FR 2741261 A		23-05-1997
			JP 9183714 A		15-07-1997
			US 5830446 A		03-11-1998
EP 0834303	A	08-04-1998	DE 19640792 C		05-03-1998
			DE 19640831 C		05-03-1998
EP 0733355	A	25-09-1996	DE 19509981 A		19-09-1996
			JP 8259426 A		08-10-1996
			US 5635461 A		03-06-1997
GB 1328108	A	30-08-1973	DE 2064591 A		15-07-1971
US 3658985	A	25-04-1972	AT 302534 B		15-09-1972
			BE 753955 A		31-12-1970
			CH 531884 A		31-12-1972
			DE 2034295 A		11-02-1971
			DK 125620 B		19-03-1973
			FR 2053194 A		16-04-1971
			GB 1307644 A		21-02-1973
			IE 34412 B		30-04-1975
			JP 48017362 B		29-05-1973
			NL 7011176 A		01-02-1971
			SE 351564 B		04-12-1972
			ZA 7004430 A		23-02-1972
CA 1255603	A	13-06-1989	NONE		
US 4312855	A	26-01-1982	US 3875071 A		01-04-1975
			US 3832310 A		27-08-1974
			AR 196483 A		06-02-1974
			AT 322708 B		10-06-1975
			AU 3452671 A		19-04-1973
			BE 775158 A		01-03-1972
			CA 970909 A		15-07-1975
			CA 955390 A		01-10-1974
			DE 2155224 A		18-05-1972
			FR 2114751 A		30-06-1972
			GB 1357960 A		26-06-1974
			IE 35776 B		12-05-1976

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IB 98/01381

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4312855 A		NL 7115801 A SE 401608 B US 3703480 A US 3726815 A	18-05-1972 22-05-1974 21-11-1972 10-04-1973
EP 0336709 A	11-10-1989	US 4866152 A CA 1337524 A JP 2011662 A JP 2542077 B US 4982603 A US 4962177 A	12-09-1989 07-11-1995 16-01-1990 09-10-1996 08-01-1991 09-10-1990
DE 2632810 A	26-01-1978	NONE	